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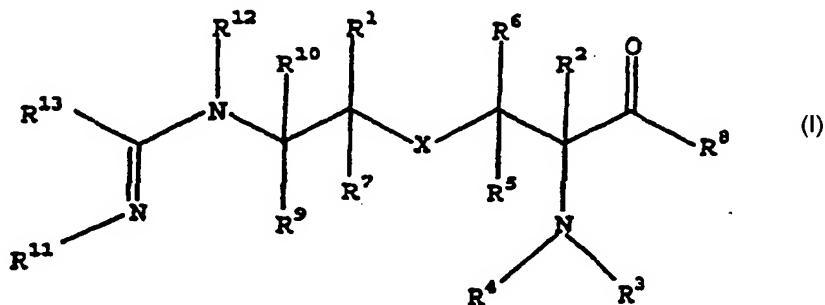
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(54) Title: AMIDINO COMPOUNDS USEFUL AS NITRIC OXIDE SYNTHASE INHIBITORS

WO 01/72702 A2



(57) Abstract: The present invention relates to amidino compounds of formula (I) in which the variables are as defined in the claims, and salts thereof. In another embodiment the present invention also provides a use of the present compounds as nitric oxide synthase inhibitors. In a further embodiment, the present invention provides methods for making the amidino compounds and other compounds.



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Amidino Compounds Useful as Nitric Oxide Synthase Inhibitors

This application claims the benefit of priority of U.S. Provisional Patent Application No. 60/191,923 filed March 24, 2000, herein incorporated by reference.

5

BACKGROUND OF THE INVENTION

Field of the Invention

10 The present invention relates to amidino compounds and their use in therapy, in particular their use as nitric oxide synthase inhibitors.

Related Art

15 It has been known since the early 1980's that the vascular relaxation caused by acetylcholine is dependent on the vascular endothelium. The endothelium-derived relaxing factor (EDRF), now known to be nitric oxide (NO) is generated in the vascular endothelium by nitric oxide synthase (NOS). The activity of NO as a vasodilator has been known for well over 100 years. In addition, NO is the active species deriving from amylnitrite, glyceryltrinitrate and other nitrovasodilators. The identification of EDRF as NO has coincided with the discovery of a biochemical pathway by which NO is synthesized from the 20 amino acid L-arginine by the enzyme NO synthase.

Nitric oxide is an endogenous stimulator of the soluble guanylate cyclase. In addition to endothelium-dependent relaxation, NO is involved in a number of biological actions including cytotoxicity of phagocytic cells and cell-to-cell communication in the central nervous system.

25 There are at least three types of NO synthase as follows:

(i) a constitutive, Ca^{++} /calmodulin dependent enzyme, located in the endothelium, that releases NO in response to receptor or physical stimulation.

(ii) a constitutive, Ca^{++} /calmodulin dependent enzyme, located in the brain, that releases NO in response to receptor or physical stimulation.

30 (iii) a Ca^{++} independent enzyme which is induced after activation of vascular smooth muscle, macrophages, endothelial cells, and a number of other cells by endotoxin and

cytokines. Once expressed this inducible nitric oxide synthase (hereinafter "iNOS") generates NO continuously for long periods.

The NO released by each of the two constitutive enzymes acts as a transduction mechanism underlying several physiological responses. The NO produced by the inducible 5 enzyme is a cytotoxic molecule for tumor cells and invading microorganisms. It also appears that adverse effects of excess NO production, in particular pathological vasodilation and tissue damage, may result largely from the NO synthesized by iNOS.

There is a growing body of evidence that NO may be involved in the degeneration of cartilage which takes place as a result of certain conditions such as arthritis and it is also 10 known that NO synthesis is increased in rheumatoid arthritis and in osteoarthritis.

Some of the NO synthase inhibitors proposed for therapeutic use are non-selective; they inhibit both the constitutive and the inducible NO synthases. Use of such a non-selective NO synthase inhibitor requires that great care be taken in order to avoid the potentially serious consequences of over-inhibition of the constitutive NO-synthase 15 including hypertension and possible thrombosis and tissue damage. In particular, in the case of the therapeutic use of L-NMMA for the treatment of toxic shock it has been recommended that the patient must be subject to continuous blood pressure monitoring throughout the treatment. Thus, while non-selective NO synthase inhibitors have therapeutic utility provided that appropriate precautions are taken, NO synthase inhibitors which are 20 selective in the sense that they inhibit the inducible NO synthase to a considerably greater extent than the constitutive isoforms of NO synthase would be of even greater therapeutic benefit and easier to use (S. Moncada and E. Higgs, FASEB J., 9, 1319-1330, 1995).

The following individual publications disclose compounds that inhibit nitric oxide synthesis and preferentially inhibit the inducible isoform of nitric oxide synthase:

- 25 PCT Patent Application No. WO 96/35677.
- PCT Patent Application No. WO 96/33175.
- PCT Patent Application No. WO 96/15120.
- PCT Patent Application No. WO 95/11014.
- PCT Patent Application No. WO 95/11231.
- 30 PCT Patent Application No. WO 99/46240.
- PCT Patent Application No. WO 95/24382.

PCT Patent Application No. WO 94/12165.
PCT Patent Application No. WO 94/14780.
PCT Patent Application No. WO 93/13055.
PCT Patent Application No. WO 99/62875.

5 European Patent No. EP0446699A1.
U.S. Patent No. 5,132,453.
U.S. Patent No. 5,684,008.
U.S. Patent No. 5,830,917.
U.S. Patent No. 5,854,251.

10 U.S. Patent No. 5,863,931.
U.S. Patent No. 5,919,787.
U.S. Patent No. 5,945,408.
U.S. Patent No. 5,981,511.

PCT Patent Application No. WO 95/25717 discloses certain amidino derivatives as
15 being useful in inhibiting inducible nitric oxide synthase.

PCT Patent Application No. WO 99/62875 discloses further amidino compounds as
being useful in inhibiting inducible nitric oxide synthase.

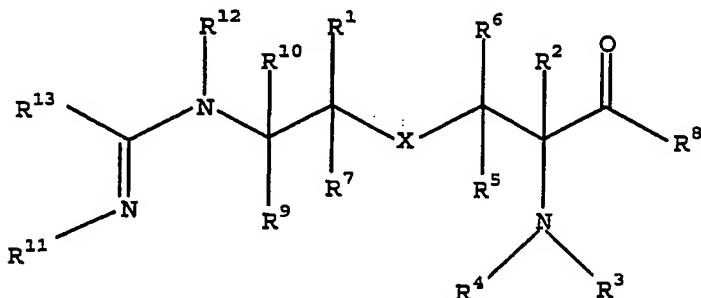
SUMMARY OF THE INVENTION

20 Compounds have now been found which have the advantage of being very efficacious as iNOS inhibitors in the human cartilage explant assay, a model for osteoarthritis. At the same time the compounds of the present invention are surprisingly unable to penetrate certain non-target organs in test systems, especially in comparison to the compounds of WO 95/25717. This surprising differentiation in expected access between the
25 target organ (cartilage) and other organs is an unexpected advantage for the compounds of the present invention.

In a broad aspect, the present invention is directed to novel compounds, pharmaceutical compositions and methods of using said compounds and compositions for inhibiting or modulating nitric oxide synthesis in a subject in need of such inhibition or
30 modulation by administering a compound which preferentially inhibits or modulates the

inducible isoform of nitric oxide synthase over the constitutive isoforms of nitric oxide synthase. It is also another object of the present invention to lower nitric oxide levels in a subject in need of such lowering. The present compounds possess useful nitric oxide synthase inhibiting activity, and are expected to be useful in the treatment or prophylaxis of a 5 disease or condition in which the synthesis or over-synthesis of nitric oxide forms a contributory part.

In one embodiment, the present invention provides a compound or a salt thereof, the 10 compound having a structure corresponding to Formula 1:



wherein:

15 X is selected from the group consisting of -S-, -S(O)-, and -S(O)2-;

 R² is selected from the group consisting of C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₅ alkoxy-C₁ alkyl, and C₁-C₅ alkylthio-C₁ alkyl;

 with respect to R³ and R⁸:

 R⁸ is selected from the group consisting of -OR¹⁴ and -N(R¹⁵)(R¹⁶); and R³ is selected 20 from the group consisting of -H, -OH, -C(O)-R¹⁷, -C(O)-O-R¹⁸, and -C(O)-S-R¹⁹; or

 R⁸ is -N(R²⁰)-, and R³ is -C(O)-, wherein R⁸ and R³ together with the atoms to which they are attached form a ring; or

 R⁸ is -O-, and R³ is -C(R²¹)(R²²)-, wherein R⁸ and R³ together with the atoms to which they are attached form a ring;

25 if R³ is -C(R²¹)(R²²)-, then R⁴ is -C(O)-O-R²³; otherwise R⁴ is -H;

R¹, R⁵, R⁶, and R⁷ independently are selected from the group consisting of -H, halogen, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, and C₁-C₅ alkoxy-C₁ alkyl;

R⁹ and R¹⁰ independently are selected from the group consisting of -H, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, and C₁-C₅ alkoxy-C₁ alkyl;

5 with respect to R¹¹ and R¹²:

R¹¹ is selected from the group consisting of -H, -OH, -C(O)-O-R²⁴, and -C(O)-S-R²⁵; and R¹² is selected from the group consisting of -H, -OH, -C(O)-O-R²⁶, and -C(O)-S-R²⁷; or R¹¹ is -O-, and R¹² is -C(O)-, wherein R¹¹ and R¹² together with the atoms to which they are attached form a ring; or

10 R¹¹ is -C(O)-, and R¹² is -O-, wherein R¹¹ and R¹² together with the atoms to which they are attached form a ring; and

R¹³ is C₁ alkyl;

R¹⁴ is selected from the group consisting of -H and C₁-C₆ alkyl; wherein when R¹⁴ is C₁-C₆ alkyl, R¹⁴ is optionally substituted by one or more moieties selected from the group 15 consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl;

with respect to R¹⁵ and R¹⁶:

R¹⁵ is selected from the group consisting of -H, alkyl, and alkoxy; and R¹⁶ is selected from the group consisting of -H, -OH, alkyl, alkoxy, -C(O)-R^{27a}, -C(O)-O-R²⁸, and -C(O)-S-R²⁹; wherein when R¹⁵ and R¹⁶ independently are alkyl or alkoxy, R¹⁵ and R¹⁶ independently 20 are optionally substituted with one or more moieties selected from the group consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl; or

R¹⁵ is -H; and R¹⁶ is selected from the group consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl;

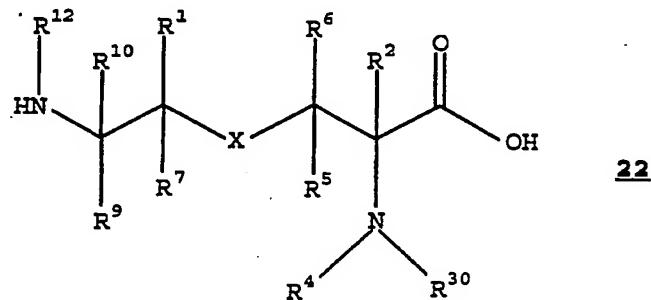
R¹⁷, R¹⁸, R¹⁹, R²⁰, R²¹, R²², R²³, R²⁴, R²⁵, R²⁶, R²⁷, R^{27a}, R²⁸, and R²⁹ independently are 25 selected from the group consisting of -H and alkyl, which is optionally substituted by one or more moieties selected from the group consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl; and

when any of R¹, R², R³, R⁴, R⁵, R⁶, R⁷, R⁸, R⁹, R¹⁰, R¹¹, R¹², R¹³, R¹⁴, R¹⁵, R¹⁶, R¹⁷, R¹⁸, R¹⁹, R²⁰, R²¹, R²², R²³, R²⁴, R²⁵, R²⁶, R²⁷, R^{27a}, R²⁸, and R²⁹ independently is a moiety selected

from the group consisting of alkyl, alkenyl, alkynyl, alkoxy, alkylthio, cycloalkyl, heterocyclyl, aryl, and heteroaryl, then the moiety is optionally substituted by one or more substituent selected from the group consisting of -OH, alkoxy, and halogen.

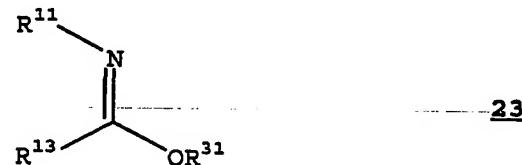
Another embodiment provides a method for the treatment or prevention of an inflammation-related disorder wherein the method comprises treating a subject in need thereof with an inflammation-related disorder treating or preventing amount of a compound of the present invention.

In yet another embodiment the present invention provides a method for the preparation of a compound of Formula 1, wherein the method comprises treating a diamine compound having a structure corresponding to Formula 22:



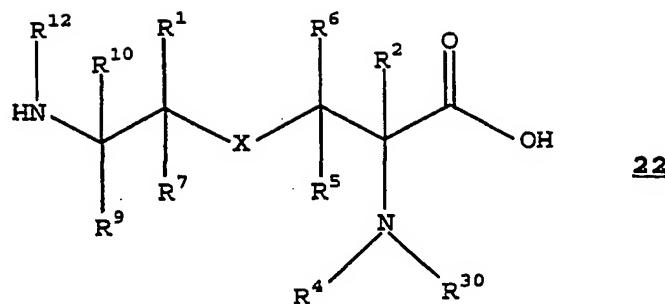
or a salt thereof, with an alkyl acetimidate having a structure corresponding to Formula 23:

15

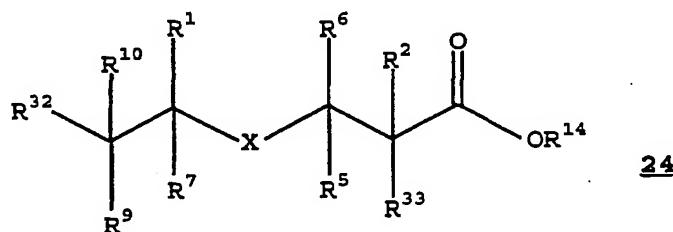


wherein R³¹ is C₁-C₆ alkyl. The treating can, if desired, be performed in the presence of an acid or a base, preferably in the presence of a base.

20 In a further embodiment of the present invention provides a method for the preparation of a diamine compound having a structure corresponding to Formula 22:



or a salt thereof, wherein R^{30} is selected from the group consisting of -H, -OH, -C(O)-R¹⁷, -C(O)-O-R¹⁸, and -C(O)-S-R¹⁹, and the other substituents are as defined above, wherein the
 5 method comprises treating a protected diamine compound having the structure corresponding to Formula 24:



10 or a salt thereof, wherein R^{33} is selected from the group consisting of -H and a protected amino group; and R^{32} is a protected amino group; and R^{14} is selected from the group consisting of -H and C₁-C₆ alkyl; wherein when R^{14} is C₁-C₆ alkyl, R^{14} is optionally substituted by one or more moieties selected from the group consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl; wherein the treating is performed with a deprotecting
 15 reagent, thereby producing the diamine compound.

DETAILED DESCRIPTION OF THE INVENTION

20 Compounds of Formula 1 will be useful for treating, among other things, inflammation in a subject, or for treating other nitric oxide synthase-mediated disorders,

such as, as an analgesic in the treatment of pain and headaches, or as an antipyretic for the treatment of fever. For example, compounds of the present invention will be useful to treat arthritis, including but not limited to rheumatoid arthritis, spondyloarthropathies, gouty arthritis, osteoarthritis, systemic lupus erythematosus, juvenile arthritis, acute rheumatic

5 arthritis, enteropathic arthritis, neuropathic arthritis, psoriatic arthritis, and pyogenic arthritis. Conditions in which the compounds of the present invention will provide an advantage in inhibiting NO production from L-arginine include arthritic conditions.

Compounds of the invention will be further useful in the treatment of asthma, bronchitis, menstrual cramps (e.g., dysmenorrhea), premature labor, tendinitis, bursitis,

10 skin-related conditions such as psoriasis, eczema, burns, sunburn, dermatitis, pancreatitis, hepatitis, and from post-operative inflammation including from ophthalmic surgery such as cataract surgery and refractive surgery. Compounds of the invention also would be useful to treat gastrointestinal conditions such as inflammatory bowel disease, Crohn's disease, gastritis, irritable bowel syndrome and ulcerative

15 colitis. Compounds of the invention would be useful for the prevention or treatment of cancer, such as colorectal cancer, and cancer of the breast, lung, prostate, bladder, cervix and skin. Compounds of the invention would be useful in treating inflammation and tissue damage in such diseases as vascular diseases, migraine headaches, periarthritis nodosa, thyroiditis, aplastic anemia, Hodgkin's disease, sclerodema,

20 rheumatic fever, type I diabetes, neuromuscular junction disease including myasthenia gravis, white matter disease including multiple sclerosis, sarcoidosis, nephrotic syndrome, Behcet's syndrome, polymyositis, gingivitis, nephritis, hypersensitivity, swelling occurring after injury, myocardial ischemia, and the like. The compounds would also be useful in the treatment of ophthalmic diseases, such as glaucoma,

25 retinitis, retinopathies, uveitis, ocular photophobia, and of inflammation and pain associated with acute injury to the eye tissue. Of particular interest among the uses of the present inventive compounds is the treatment of glaucoma, especially where symptoms of glaucoma are caused by the production of nitric oxide, such as in nitric oxide-mediated nerve damage. The compounds would also be useful in the treatment

30 of pulmonary inflammation, such as that associated with viral infections and cystic fibrosis. The compounds would also be useful for the treatment of certain central

nervous system disorders, such as cortical dementias including Alzheimer's disease, and central nervous system damage resulting from stroke, ischemia and trauma. The compounds of the invention are useful as anti-inflammatory agents, such as for the treatment of arthritis, with the additional benefit of having significantly less harmful side effects. These compounds would also be useful in the treatment of allergic rhinitis, respiratory distress syndrome, endotoxin shock syndrome, and atherosclerosis. The compounds would also be useful in the treatment of pain, but not limited to postoperative pain, dental pain, muscular pain, and pain resulting from cancer. The compounds would be useful for the prevention of dementias, such as Alzheimer's disease.

Besides being useful for human treatment, these compounds are also useful for veterinary treatment of companion animals, exotic animals and farm animals, including mammals, rodents, and the like. More preferred animals include horses, dogs, and cats.

The present compounds may also be used in co-therapies, partially or completely, in place of other conventional antiinflammatory therapies, such as together with steroids, NSAIDs, COX-2 selective inhibitors, 5-lipoxygenase inhibitors, LTB₄ antagonists and LTA₄ hydrolase inhibitors.

Other conditions in which the compounds of the present invention will provide an advantage in inhibiting NO inhibition include cardiovascular ischemia, diabetes (type I or type II), congestive heart failure, myocarditis, atherosclerosis, migraine, glaucoma, aortic aneurysm, reflux esophagitis, diarrhea, irritable bowel syndrome, cystic fibrosis, emphysema, asthma, bronchiectasis, hyperalgesia (allodynia), cerebral ischemia (both focal ischemia, thrombotic stroke and global ischemia (for example, secondary to cardiac arrest), multiple sclerosis and other central nervous system disorders mediated by NO, for example Parkinson's disease. Further neurodegenerative disorders in which NO inhibition may be useful include nerve degeneration or nerve necrosis in disorders such as hypoxia, hypoglycemia, epilepsy, and in cases of central nervous system (CNS) trauma (such as spinal cord and head injury), hyperbaric oxygen convulsions and toxicity, dementia e.g. pre-senile dementia, and AIDS-related dementia, cachexia, Sydenham's chorea, Huntington's disease, Amyotrophic Lateral Sclerosis, Korsakoff's disease, imbecility relating to a cerebral vessel

disorder, sleeping disorders, schizophrenia, depression, depression or other symptoms associated with Premenstrual Syndrome (PMS), anxiety and septic shock.

The compounds of the present invention will also be useful in the treatment of pain including somatogenic (either nociceptive or neuropathic), both acute and chronic. A nitric oxide inhibitor could be used in any situation including neuropathic pain that a common NSAID or opioid analgesic would traditionally be administered.

Still other disorders or conditions which will be advantageously treated by the compounds of the present invention include treatment of prevention of opiate tolerance in patients needing protracted opiate analgesics, and benzodiazepine tolerance in patients taking benzodiazepines, and other addictive behavior, for example, nicotine addiction, alcoholism, and eating disorders. The compounds and methods of the present invention will also be useful in the treatment or prevention of drug withdrawal symptoms, for example treatment or prevention of symptoms of withdrawal from opiate, alcohol, or tobacco addiction. The present inventive compounds may also be useful to prevent tissue damage when therapeutically combined with antibacterial or antiviral agents.

The compounds of the present invention will also be useful in inhibiting NO production from L-arginine including systemic hypotension associated with septic and/or toxic hemorrhagic shock induced by a wide variety of agents; therapy with cytokines such as TNF, IL-1 and IL-2; and as an adjuvant to short term immunosuppression in transplant therapy.

The present invention is further directed to the use of the compounds of the present invention for the treatment and prevention of neoplasias. The neoplasias that will be treatable or preventable by the compounds and methods of the present invention include brain cancer, bone cancer, a leukemia, a lymphoma, epithelial cell-derived neoplasia (epithelial carcinoma) such as basal cell carcinoma, adenocarcinoma, gastrointestinal cancer such as lip cancer, mouth cancer, esophageal cancer, small bowel cancer and stomach cancer, colon cancer, liver cancer, bladder cancer, pancreas cancer, ovary cancer, cervical cancer, lung cancer, breast cancer and skin cancer, such as squamous cell and basal cell cancers, prostate cancer, renal cell carcinoma, and other known cancers that effect epithelial cells throughout the body. Preferably, the neoplasia is selected from gastrointestinal cancer, liver cancer, bladder cancer, pancreas cancer, ovary cancer, prostate cancer, cervical cancer,

lung cancer, breast cancer and skin cancer, such as squamous cell and basal cell cancers. The present compounds and methods can also be used to treat the fibrosis which occurs with radiation therapy. The present compounds and methods can be used to treat subjects having adenomatous polyps, including those with familial adenomatous polyposis (FAP).

5 Additionally, the present compounds and methods can be used to prevent polyps from forming in patients at risk of FAP.

Conjunctive treatment of a compound of the present invention with another antineoplastic agent will produce a synergistic effect or alternatively reduce the toxic side effects associated with chemotherapy by reducing the therapeutic dose of the side effect-causing agent needed for therapeutic efficacy or by directly reducing symptoms of toxic side effects caused by the side effect-causing agent. A compound of the present invention will further be useful as an adjunct to radiation therapy to reduce side effects or enhance efficacy.

In the present invention, another agent which can be combined therapeutically with a compound of the present invention includes any therapeutic agent which is capable of

10 inhibiting the enzyme cyclooxygenase-2 (“COX-2”). Preferably such COX-2 inhibiting agents inhibit COX-2 selectively relative to the enzyme cyclooxygenase-1 (“COX-1”). Such a COX-2 inhibitor is known as a “COX-2 selective inhibitor”. More preferably, a compound of the present invention can be therapeutically combined with a COX-2 selective inhibitor wherein the COX-2 selective inhibitor selectively inhibits COX-2 at a ratio of at least 10:1

15 relative to inhibition of COX-1, more preferably at least 30:1, and still more preferably at least 50:1 in an *in vitro* test. COX-2 selective inhibitors useful in therapeutic combination with the compounds of the present invention include celecoxib, valdecoxib, deracoxib, etoricoxib, rofecoxib, ABT-963 (2-(3,4-difluorophenyl)-4-(3-hydroxy-3-methyl-1-butoxy)-5-[4-(methylsulfonyl)phenyl-3(2H)-pyridazinone; described in PCT Patent Application No.

20 20 WO 00/24719), or meloxicam. A compound of the present invention can also be advantageously used in therapeutic combination with a prodrug of a COX-2 selective inhibitor, for example parecoxib.

Another chemotherapeutic agent which will be useful in combination with a compound of the present invention can be selected, for example, from the following non-comprehensive and non-limiting list:

Alpha-difluoromethylornithine (DFMO), 5-FU-fibrinogen, acanthifolic acid, aminothiadiazole, brequinar sodium, carmofur, Ciba-Geigy CGP-30694, cyclopentyl cytosine, cytarabine phosphate stearate, cytarabine conjugates, Lilly DATHF, Merrel Dow DDFC, dezaguanine, dideoxycytidine, dideoxyguanosine, didox, Yoshitomi DMDC,

5 doxifluridine, Wellcome EHNA, Merck & Co. EX-015, fazarabine, flouxuridine, fludarabine phosphate, 5-fluorouracil, N-(2'-furanidyl)-5-fluorouracil, Daiichi Seiyaku FO-152, isopropyl pyrrolizine, Lilly LY-188011, Lilly LY-264618, methobenzaprim, methotrexate, Wellcome MZPES, norspermidine, NCI NSC-127716, NCI NSC-264880, NCI NSC-39661, NCI NSC-612567, Warner-Lambert PALA, pentostatin, piritrexim, plicamycin, Asahi

10 Chemical PL-AC, Takeda TAC-788, thioguanine, tiazofurin, Erbamont TIF, trimetrexate, tyrosine kinase inhibitors, tyrosine protein kinase inhibitors, Taiho UFT, uricytin, Shionogi 254-S, aldo-phosphamide analogues, altretamine, anaxirone, Boehringer Mannheim BBR-2207, bestrabucil, budotitane, Wakunaga CA-102, carboplatin, carmustine, Chinoin-139, Chinoin-153, chlorambucil, cisplatin, cyclophosphamide, American Cyanamid CL-286558,

15 Sanofi CY-233, cyplatate, Degussa D-19-384, Sumimoto DACHP(Myr)2, diphenylspiromustine, diplatinum cytostatic, Erba distamycin derivatives, Chugai DWA-2114R, ITI E09, elmustine, Erbamont FCE-24517, estramustine phosphate sodium, fotemustine, Unimed G-6-M, Chinoin GYKI-17230, hepsul-fam, ifosfamide, iproplatin, lomustine, mafosfamide, mitolactol, Nippon Kayaku NK-121, NCI NSC-264395, NCI NSC-

20 342215, oxaliplatin, Upjohn PCNU, prednimustine, Proter PTT-119, ranimustine, semustine, SmithKline SK&F-101772, Yakult Honsha SN-22, spiromus-tine, Tanabe Seiyaku TA-077, tauromustine, temozolomide, teroxirone, tetraplatin, trimelamol, Taiho 4181-A, aclarubicin, actinomycin D, actinoplanone, Erbamont ADR-456, aeroplysinin derivative, Ajinomoto AN-201-II, Ajinomoto AN-3, Nippon Soda anisomycins, anthracycline, azino-mycin-A,

25 bisucaberin, Bristol-Myers BL-6859, Bristol-Myers BMY-25067, Bristol-Myers BMY-25551, Bristol-Myers BMY-26605, Bristol-Myers BMY-27557, Bristol-Myers BMY-28438, bleomycin sulfate, bryostatin-1, Taiho C-1027, calichemycin, chromoximycin, dactinomycin, daunorubicin, Kyowa Hakko DC-102, Kyowa Hakko DC-79, Kyowa Hakko DC-88A, Kyowa Hakko DC89-A1, Kyowa Hakko DC92-B, ditrisarubicin B, Shionogi

30 DOB-41, doxorubicin, doxorubicin-fibrinogen, elsamicin-A, epirubicin, erbstatin, esorubicin, esperamicin-A1, esperamicin-Alb, Erbamont FCE-21954, Fujisawa FK-973,

fosfriecin, Fujisawa FR-900482, glidobactin, gregatin-A, grincamycin, herbimycin, idarubicin, illudins, kazusamycin, kesarirhodins, Kyowa Hakko KM-5539, Kirin Brewery KRN-8602, Kyowa Hakko KT-5432, Kyowa Hakko KT-5594, Kyowa Hakko KT-6149, American Cyanamid LL-D49194, Meiji Seika ME 2303, menogaril, mitomycin,

5 mitoxantrone, SmithKline M-TAG, neoenactin, Nippon Kayaku NK-313, Nippon Kayaku NKT-01, SRI International NSC-357704, oxalysine, oxaunomycin, peplomycin, pilatin, pirarubicin, porothramycin, pyrindamycin A, Tobishi RA-I, rapamycin, rhizoxin, rodorubicin, sibanomicin, siwenmycin, Sumitomo SM-5887, Snow Brand SN-706, Snow Brand SN-07, sorangicin-A, sparsomycin, SS Pharmaceutical SS-21020, SS Pharmaceutical

10 SS-7313B, SS Pharmaceutical SS-9816B, steffimycin B, Taiho 4181-2, talisomycin, Takeda TAN-868A, terpentecin, thrazine, tricrozarin A, Upjohn U-73975, Kyowa Hakko UCN-10028A, Fujisawa WF-3405, Yoshitomi Y-25024 zorubicin, alpha-carotene, alpha-difluoromethyl-arginine, acitretin, Biotec AD-5, Kyorin AHC-52, alstonine, amonafide, amphethinile, amsacrine, Angiostat, ankinomycin, anti-neoplaston A10, antineoplaston A2,

15 antineoplaston A3, antineoplaston A5, antineoplaston AS2-1, Henkel APD, aphidicolin glycinate, asparaginase, Avarol, baccharin, batracylin, benfluron, benzotript, Ipsen-Beaufour BIM-23015, bisantrene, Bristo-Myers BMY-40481, Vestar boron-10, bromofosfamide, Wellcome BW-502, Wellcome BW-773, caracemide, carmethylizole hydrochloride, Ajinomoto CDAF, chlorsulfaquinoxalone, Chemex CHX-2053, Chemex CHX-100, Warner-

20 Lambert CI-921, Warner-Lambert CI-937, Warner-Lambert CI-941, Warner-Lambert CI-958, clanfenur, claviridenone, ICN compound 1259, ICN compound 4711, Contracan, Yakult Honsha CPT-11, crisnatol, curaderm, cytochalasin B, cytarabine, cytocytin, Merz D-609, DABIS maleate, dacarbazine, datelliptinium, didemnin-B, dihaematoporphyrin ether, dihydrolenperone, dinaline, distamycin, Toyo Pharmar DM-341, Toyo Pharmar DM-75,

25 Daiichi Seiyaku DN-9693, elliprabin, elliptinium acetate, Tsumura EPMTC, ergotamine, etoposide, etretinate, fenretinide, Fujisawa FR-57704, gallium nitrate, genkwadaphnin, Chugai GLA-43, Glaxo GR-63178, grifolan NMF-5N, hexadecylphosphocholine, Green Cross HO-221, homoharringtonine, hydroxyurea, BTG ICRF-187, ilmofosine, isoglutamine, isotretinoin, Otsuka JI-36, Ramot K-477, Otsuak K-76COONa, Kureha Chemical K-AM,

30 MECT Corp KI-8110, American Cyanamid L-623, leukoregulin, lonidamine, Lundbeck LU-23-112, Lilly LY-186641, NCI (US) MAP, marycin, Merrel Dow MDL-27048, Medco

MEDR-340, merbarone, merocyanine derivatives, methylanilinoacridine, Molecular Genetics MGI-136, minactivin, mitonafide, mitoquidone, mopidamol, motretinide, Zenyaku Kogyo MST-16, N-(retinoyl)amino acids, Nissrin Flour Milling N-021, N-acylated-dehydroalanines, nafazatrom, Taisho NCU-190, nocodazole derivative, Normosang, NCI

5 NSC-145813, NCI NSC-361456, NCI NSC-604782, NCI NSC-95580, octreotide, Ono ONO-112, oquizanocene, Akzo Org-10172, pancratistatin, pazelliptine, Warner-Lambert PD-111707, Warner-Lambert PD-115934, Warner-Lambert PD-131141, Pierre Fabre PE-1001, ICRT peptide D, piroxantrone, polyhaematoxyporphyrin, polypreic acid, Efamol porphyrin, probimane, procarbazine, proglumide, Invitron protease nexin I, Tobishi RA-700, razoxane,

10 Sapporo Breweries RBS, restrictin-P, retelliptine, retinoic acid, Rhone-Poulenc RP-49532, Rhone-Poulenc RP-56976, SmithKline SK&F-104864, Sumitomo SM-108, Kuraray SMANCS, SeaPharm SP-10094, spatal, spirocyclopropane derivatives, spirogermanium, Unimed, SS Pharmaceutical SS-554, strypoldinone, Stypoldione, Suntory SUN 0237, Suntory SUN 2071, superoxide dismutase, Toyama T-506, Toyama T-680, taxol, Teijin TEI-

15 0303, teniposide, thaliblastine, Eastman Kodak TJB-29, tocotrienol, Topostin, Teijin TT-82, Kyowa Hakko UCN-01, Kyowa Hakko UCN-1028, ukrain, Eastman Kodak USB-006, vinblastine sulfate, vincristine, vindesine, vinestramide, vinorelbine, vinriptol, vinzolidine, withanolides, Yamanouchi YM-534, uroguanylin, combretastatin, dolastatin, idarubicin, epirubicin, estramustine, cyclophosphamide, 9-amino-2-(S)-camptothecin, topotecan,

20 irinotecan (Camptosar), exemestane, decapeptyl (tryptorelin), or an omega-3 fatty acid.

Examples of radioprotective agents which may be used in a combination therapy with the compounds of this invention include AD-5, adchnon, amifostine analogues, detox, dimesna, I-102, MM-159, N-acylated-dehydroalanines, TGF- Genentech, tiprotimod, amifostine, WR-151327, FUT-187, ketoprofen transdermal, nabumetone, superoxide dismutase (Chiron) and superoxide dismutase Enzon.

The compounds of the present invention will also be useful in treatment or prevention of angiogenesis-related disorders or conditions, for example, tumor growth, metastasis, macular degeneration, and atherosclerosis.

In a further embodiment, the present invention also provides therapeutic combinations for the treatment or prevention of ophthalmic disorders or conditions such as glaucoma. For example the present inventive compounds advantageously will be used in

therapeutic combination with a drug which reduces the intraocular pressure of patients afflicted with glaucoma. Such intraocular pressure-reducing drugs include without limitation latanoprost, travoprost, bimatoprost, or unoprostone. The therapeutic combination of a compound of the present invention plus an intraocular pressure-reducing drug will be useful 5 because each is believed to achieve its effects by affecting a different mechanism.

In another combination of the present invention, the present inventive compounds can be used in therapeutic combination with an antihyperlipidemic or cholesterol-lowering drug such as a benzothiepine or a benzothiazepine antihyperlipidemic drug. Examples of benzothiepine antihyperlipidemic drugs useful in the present inventive therapeutic 10 combination can be found in U.S. Patent No. 5,994,391, herein incorporated by reference. Some benzothiazepine antihyperlipidemic drugs are described in WO 93/16055. Alternatively, the antihyperlipidemic or cholesterol-lowering drug useful in combination with a compound of the present invention can be an HMG Co-A reductase inhibitor. Examples of HMG Co-A reductase inhibitors useful in the present therapeutic combination 15 include, individually, benfluorex, fluvastatin, lovastatin, provastatin, simvastatin, atorvastatin, cerivastatin, bervastatin, ZD-9720 (described in PCT Patent Application No. WO 97/06802), ZD-4522 (CAS No. 147098-20-2 for the calcium salt; CAS No. 147098-18-8 for the sodium salt; described in European Patent No. EP 521471), BMS 180431 (CAS No. 129829-03-4), or NK-104 (CAS No. 141750-63-2). The therapeutic combination of a 20 compound of the present invention plus an antihyperlipidemic or cholesterol-lowering drug will be useful, for example, in reducing the risk of formation of atherosclerotic lesions in blood vessels. For example, atherosclerotic lesions often initiate at inflamed sites in blood vessels. It is established that antihyperlipidemic or cholesterol-lowering drug reduce risk of formation of atherosclerotic lesions by lowering lipid levels in blood. Without limiting the 25 invention to a single mechanism of action, it is believed that one way the compounds of the present combination will work in concert to provide improved control of atherosclerotic lesions by, for example, reducing inflammation of the blood vessels in concert with lowering blood lipid levels.

In another embodiment of the invention, the present compounds can be used in 30 combination with other compounds or therapies for the treatment of central nervous conditions or disorders such as migraine. For example, the present compounds can be used

in therapeutic combination with caffeine, a 5-HT-1B/1D agonist (for example, a triptan such as sumatriptan, naratriptan, zolmitriptan, rizatriptan, almotriptan, or frovatriptan), a dopamine D4 antagonist (e.g., sonepiprazole), aspirin, acetaminophen, ibuprofen, indomethacin, naproxen sodium, isometheptene, dichloralphenazone, butalbital, an ergot 5 alkaloid (e.g., ergotamine, dihydroergotamine, bromocriptine, ergonovine, or methyl ergonovine), a tricyclic antidepressant (e.g., amitriptyline or nortriptyline), a serotonergic antagonist (e.g., methysergide or cyproheptadine), a beta-andrenergic antagonist (e.g., propranolol, timolol, atenolol, nadolol, or metprolol), or a monoamine oxidase inhibitor (e.g., phenelzine or isocarboxazid).

10 A further embodiment provides a therapeutic combination of a compound of the present invention with an opioid compound. Opioid compounds useful in this combination include without limitation morphine, methadone, hydromorphone, oxymorphone, levorphanol, levallorphan, codeine, dihydrocodeine, dihydrohydroxycodeinone, pentazocine, hydrocodone, oxycodone, nalmefene, etorphine, levorphanol, fentanyl, sufentanil, DAMGO, 15 butorphanol, buprenorphine, naloxone, naltrexone, CTOP, diprenorphine, beta-funaltrexamine, naloxonazine, nalorphine, pentazocine, nalbuphine, naloxone benzoylhydrazone, bremazocine, ethylketocyclazocine, U50,488, U69,593, spiradoline, norbinaltorphimine, naltrindole, DPDPE, [D- α^2 , glu 4]deltorphin, DSLET, met-enkephalin, leu-enkaphalin, beta-endorphin, dynorphin A, dynorphin B, and alpha-neoendorphin. An 20 advantage to the combination of the present invention with an opioid compound is that the present inventive compounds will allow a reduction in the dose of the opioid compound, thereby reducing the risk or severity of opioid side effects, such as opioid addiction.

The term "alkyl", alone or in combination, means an acyclic alkyl radical, linear or branched, preferably containing from 1 to about 10 carbon atoms and more preferably 25 containing from 1 to about 6 carbon atoms. "Alkyl" also encompasses cyclic alkyl radicals containing from 3 to about 7 carbon atoms, preferably from 3 to 5 carbon atoms. Said alkyl radicals can be optionally substituted with groups as defined below. Examples of such radicals include methyl, ethyl, chloroethyl, hydroxyethyl, n-propyl, isopropyl, n-butyl, cyanobutyl, isobutyl, sec-butyl, tert-butyl, pentyl, aminopentyl, iso-amyl, hexyl, octyl and 30 the like.

The term "alkenyl" refers to an unsaturated, acyclic hydrocarbon radical, linear or branched, in so much as it contains at least one double bond. Such radicals containing from 2 to about 6 carbon atoms, preferably from 2 to about 4 carbon atoms, more preferably from 2 to about 3 carbon atoms. Said alkenyl radicals may be optionally substituted with groups as defined below. Examples of suitable alkenyl radicals include propenyl, 2-chloropropenyl, buten-1-yl, isobutenyl, penten-1-yl, 2-methylbuten-1-yl, 3-methylbuten-1-yl, hexen-1-yl, 3-hydroxyhexen-1-yl, hepten-1-yl, and octen-1-yl, and the like.

The term "alkynyl" refers to an unsaturated, acyclic hydrocarbon radical, linear or branched, in so much as it contains one or more triple bonds, such radicals containing 2 to 10 about 6 carbon atoms, preferably from 2 to about 4 carbon atoms, more preferably from 2 to about 3 carbon atoms. Said alkynyl radicals may be optionally substituted with groups as defined below. Examples of suitable alkynyl radicals include ethynyl, propynyl, hydroxypropynyl, butyn-1-yl, butyn-2-yl, pentyn-1-yl, pentyn-2-yl, 4-methoxypentyn-2-yl, 3-methylbutyn-1-yl, hexyn-1-yl, hexyn-2-yl, hexyn-3-yl, 3,3-dimethylbutyn-1-yl radicals and the like.

The term "alkoxy" embrace linear or branched oxy-containing radicals each having alkyl portions of 1 to about 6 carbon atoms, preferably 1 to about 3 carbon atoms, such as a methoxy radical. The term "alkoxyalkyl" also embraces alkyl radicals having one or more alkoxy radicals attached to the alkyl radical, that is, to form monoalkoxyalkyl and dialkoxyalkyl radicals. Examples of such radicals include methoxy, ethoxy, propoxy, butoxy and *tert*-butoxy alkyls. The "alkoxy" radicals may be further substituted with one or more halo atoms, such as fluoro, chloro or bromo, to provide "haloalkoxy" radicals. Examples of such radicals include fluoromethoxy, chloromethoxy, trifluoromethoxy, difluoromethoxy, trifluoroethoxy, fluoroethoxy, tetrafluoroethoxy, pentafluoroethoxy, and fluoropropoxy.

The term "alkylthio" embraces radicals containing a linear or branched alkyl radical, of 1 to about 6 carbon atoms, attached to a divalent sulfur atom. An example of "lower alkylthio" is methylthio ($\text{CH}_3\text{-S-}$).

The term "alkylthioalkyl" embraces alkylthio radicals, attached to an alkyl group. Examples of such radicals include methylthiomethyl.

The term "halo" means halogens such as fluorine, chlorine, bromine or iodine atoms.

The term "heterocyclyl" means a saturated or unsaturated mono- or multi-ring carbocycle wherein one or more carbon atoms is replaced by N, S, P, or O. This includes, for example, the following structures:

5



wherein Z, Z¹, Z² or Z³ is C, S, P, O, or N, with the proviso that one of Z, Z¹, Z² or Z³ is other than carbon, but is not O or S when attached to another Z atom by a double bond or 10 when attached to another O or S atom. Furthermore, the optional substituents are understood to be attached to Z, Z¹, Z² or Z³ only when each is C. The term "heterocyclyl" also includes fully saturated ring structures such as piperazinyl, dioxanyl, tetrahydrofuranyl, oxiranyl, aziridinyl, morpholinyl, pyrrolidinyl, piperidinyl, thiazolidinyl, and others. The term "heterocyclyl" also includes partially unsaturated ring structures such as dihydrofuranyl, 15 pyrazolinyl, imidazolinyl, pyrrolinyl, chromanyl, dihydrothiophenyl, and others.

The term "heteroaryl" means a fully unsaturated heterocycle.

In either "heterocycle" or "heteroaryl," the point of attachment to the molecule of interest can be at the heteroatom or elsewhere within the ring.

The term "cycloalkyl" means a mono- or multi-ringed carbocycle wherein each ring 20 contains three to about seven carbon atoms, preferably three to about five carbon atoms. Examples include radicals such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloalkenyl, and cycloheptyl. The term "cycloalkyl" additionally encompasses spiro systems wherein the cycloalkyl ring has a carbon ring atom in common with the seven-membered heterocyclic ring of the benzothiepine.

25 The term "oxo" means a doubly bonded oxygen.

The term "alkoxy" means a radical comprising an alkyl radical that is bonded to an oxygen atom, such as a methoxy radical. More preferred alkoxy radicals are "lower alkoxy" radicals having one to about ten carbon atoms. Still more preferred alkoxy radicals have one

to about six carbon atoms. Examples of such radicals include methoxy, ethoxy, propoxy, isopropoxy, butoxy and tert-butoxy.

The term "aryl" means a fully unsaturated mono- or multi-ring carbocycle, including, but not limited to, substituted or unsubstituted phenyl, naphthyl, or anthracenyl.

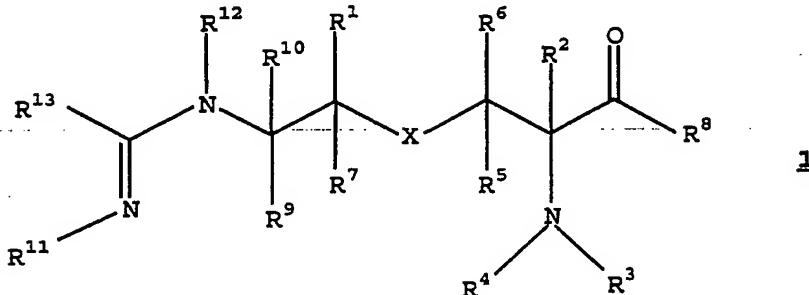
5 The term "Combination therapy" means the administration of two or more therapeutic agents to treat a therapeutic condition or disorder described in the present disclosure, for example atherosclerosis, pain, inflammation, migraine, neoplasia, angiogenesis-related condition or disorder, or other. Such administration encompasses co-administration of these therapeutic agents in a substantially simultaneous manner, such as in

10 a single capsule having a fixed ratio of active ingredients or in multiple, separate capsules for each active ingredient. In addition, such administration also encompasses use of each type of therapeutic agent in a sequential manner. In either case, the treatment regimen will provide beneficial effects of the drug combination in treating the conditions or disorders described herein.

15 The phrase "therapeutically effective" is intended to qualify the combined amount of active ingredients in the combination therapy. This combined amount will achieve the goal of reducing or eliminating the hyperlipidemic condition.

In one embodiment, the present invention provides a compound or a salt thereof, the compound having a structure corresponding to Formula 1:

20



In the structure of Formula 1, X is selected from the group consisting of -S-, -S(O)-, and -S(O)₂-.

25 Preferably, X is -S-. R² is selected from the group consisting of C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₅ alkoxy-C₁ alkyl, and C₁-C₅ alkylthio-C₁ alkyl wherein

each of these groups is optionally substituted by one or more substituent selected from the group consisting of -OH, alkoxy, and halogen. Preferably, R² is C₁-C₆ alkyl optionally substituted with a substituent selected from the group consisting of -OH, alkoxy, and halogen. With respect to R³ and R⁸, R⁸ is selected from the group consisting of -OR¹⁴ and -

5 N(R¹⁵)(R¹⁶), and R³ is selected from the group consisting of -H, -OH, -C(O)-R¹⁷, -C(O)-O- R¹⁸, and -C(O)-S-R¹⁹; or R⁸ is -N(R²⁰)-, and R³ is -C(O)-, wherein R⁸ and R³ together with the atoms to which they are attached form a ring; or R⁸ is -O-, and R³ is -C(R²¹)(R²²)-, wherein R⁸ and R³ together with the atoms to which they are attached form a ring. If R³ is -

10 C(R²¹)(R²²)-, then R⁴ is -C(O)-O-R²³; otherwise R⁴ is -H. R¹, R⁵, R⁶, and R⁷ independently

15 are selected from the group consisting of -H, halogen, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, and C₁-C₅ alkoxy-C₁ alkyl. R⁹ and R¹⁰ independently are selected from the group consisting of -H, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, and C₁-C₅ alkoxy-C₁ alkyl. With respect to R¹¹ and R¹², R¹¹ is selected from the group consisting of -H, -OH, -C(O)-O- R²⁴, and -C(O)-S-R²⁵, and R¹² is selected from the group consisting of -H, -OH, -C(O)-O-R²⁶,

15 and -C(O)-S-R²⁷; or R¹¹ is -O-, and R¹² is -C(O)-, wherein R¹¹ and R¹² together with the atoms to which they are attached form a ring; or R¹¹ is -C(O)-, and R¹² is -O-, wherein R¹¹ and R¹² together with the atoms to which they are attached form a ring. R¹³ is C₁ alkyl. R¹⁴ is selected from the group consisting of -H and C₁-C₆ alkyl, wherein when R¹⁴ is C₁-C₆ alkyl, R¹⁴ is optionally substituted by one or more moieties selected from the group

20 consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl. With respect to R¹⁵ and R¹⁶, R¹⁵ is selected from the group consisting of -H, alkyl, and alkoxy, and R¹⁶ is selected from the group consisting of -H, -OH, alkyl, alkoxy, -C(O)-R^{27a}, -C(O)-O-R²⁸, and -C(O)-S-R²⁹; wherein when R¹⁵ and R¹⁶ independently are alkyl or alkoxy, R¹⁵ and R¹⁶ independently are optionally substituted with one or more moieties selected from the group consisting of

25 cycloalkyl, heterocyclyl, aryl, and heteroaryl; or R¹⁵ is -H; and R¹⁶ is selected from the group consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl. R¹⁷, R¹⁸, R¹⁹, R²⁰, R²¹, R²², R²³, R²⁴, R²⁵, R²⁶, R²⁷, R^{27a}, R²⁸, and R²⁹ independently are selected from the group consisting of -H and alkyl, wherein alkyl is optionally substituted by one or more moieties selected from the group consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl. When any of R¹, R²,

$R^3, R^4, R^5, R^6, R^7, R^8, R^9, R^{10}, R^{11}, R^{12}, R^{13}, R^{14}, R^{15}, R^{16}, R^{17}, R^{18}, R^{19}, R^{20}, R^{21}, R^{22}, R^{23}, R^{24}, R^{25}, R^{26}, R^{27}, R^{27a}, R^{28}$, and R^{29} independently is a moiety selected from the group consisting of alkyl, alkenyl, alkynyl, alkoxy, alkylthio, cycloalkyl, heterocyclyl, aryl, and heteroaryl, then the moiety is optionally substituted by one or more substituent selected from the group 5 consisting of -OH, alkoxy, and halogen.

In a preferred embodiment, R^8 is -OH. When R^8 is -OH, preferably X is S. In a further embodiment, R^1, R^5, R^6, R^7, R^9 , and R^{10} independently are selected from the group consisting of -H and C_1 - C_3 alkyl. Preferably $R^5, R^6, R^7, R^9, R^{10}$ each are -H. R^{13} can be a variety of groups, for example fluoromethyl or methyl. R^1 can be C_1 - C_6 alkyl optionally 10 substituted with a substituent selected from the group consisting of -OH and halogen; preferably R^1 is C_1 alkyl optionally substituted with halogen; more preferably R^1 is selected from the group consisting of fluoromethyl, hydroxymethyl, and methyl. In one important embodiment R^1 can be methyl. Alternatively, R^1 can be fluoromethyl. In another alternative R^1 can be hydroxymethyl. In another embodiment, R^2 is C_1 - C_6 alkyl optionally substituted 15 with a substituent selected from the group consisting of -OH, alkoxy, and halogen. In one preferred embodiment R^2 is C_1 alkyl optionally substituted with halogen. For example, R^2 can be methyl. Alternatively, R^2 can be fluoromethyl. In yet another example, R^2 can be hydroxymethyl. In still another example, R^2 can be methoxymethyl.

In the compounds of the present invention, it is preferred that R^3, R^4, R^{11} and R^{12} each 20 is -H. In this embodiment, it is further preferred that R^1, R^5, R^6, R^7, R^9 , and R^{10} independently are selected from the group consisting of -H and C_1 - C_3 alkyl. Preferably $R^5, R^6, R^7, R^9, R^{10}$ each is -H. In this further embodiment, R^{13} can be, for example, fluoromethyl, or in another example R^{13} can be methyl. In preferred compounds of these examples, R^2 is C_1 - C_6 alkyl optionally substituted with a substituent selected from the group consisting of - 25 OH, alkoxy, and halogen. Preferably R^2 is C_1 alkyl optionally substituted with halogen. In one such example R^2 is fluoromethyl. In another example R^2 is methyl. Alternatively R^2 can be hydroxymethyl. In another alternative, R^2 can be methoxymethyl.

When R^{13} is methyl, R^1 can be, for example, -H or C_1 - C_6 alkyl optionally substituted with a substituent selected from the group consisting of -OH and halogen. In a preferred embodiment R^1 is -H. Alternatively, R^1 can be C_1 - C_6 alkyl optionally substituted with a substituent selected from the group consisting of -OH and halogen. For example R^1 can be 5 methyl, ethyl, n-propyl, i-propyl, n-butyl, sec-butyl, isobutyl, t-butyl, a pentyl isomer, or a hexyl isomer. For example, R^1 can be ethyl. Alternatively, R^1 can be C_1 alkyl optionally substituted with a substituent selected from the group consisting of -OH and halogen; for example R^1 can be methyl. Alternatively, R^1 can be fluoromethyl. In another alternative, R^1 can be hydroxymethyl.

10 In another embodiment R^8 can be - OR^{14} . R^{14} can be as defined above. Preferably R^{14} is C_1 - C_6 alkyl optionally substituted by one or more moieties selected from the group consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl; more preferably R^{14} is C_1 - C_3 alkyl; and more preferably still R^{14} is methyl. In yet another embodiment of compound 1, R^8 can be - $N(R^{15})(R^{16})$, wherein R^{15} and R^{16} are as defined above. In still another embodiment 15 R^8 can be - $N(R^{20})-$, and R^3 can be - $C(O)-$, wherein R^8 and R^3 together with the atoms to which they are attached form a ring. In another embodiment still, R^8 can be - $O-$, and R^3 can be - $C(R^{21})(R^{22})-$, wherein R^8 and R^3 together with the atoms to which they are attached form a ring.

20 In the compound of Formula 1, R^{11} can be selected from the group consisting of -OH, - $C(O)-O-R^{24}$, and - $C(O)-S-R^{25}$. Preferably R^{11} is -OH. In a further embodiment R^{12} is -H when R^{11} is -OH.

25 However, the present invention also provides useful compounds of Formula 1 in which R^{11} is - $O-$, and R^{12} is - $C(O)-$, wherein R^{11} and R^{12} together with the atoms to which they are attached form a ring. In another useful embodiment R^{11} is - $C(O)-$, and R^{12} is - $O-$, wherein R^{11} and R^{12} together with the atoms to which they are attached form a ring. Alternatively, R^{12} can be selected from the group consisting of -OH, - $C(O)-O-R^{26}$, and - $C(O)-S-R^{27}$. In this alternative, R^{11} is preferably -H.

The present invention also provides pharmaceutically-acceptable salts of the compounds of Formula 1. For example, such a pharmaceutically acceptable salt can be one

in which the present inventive compound is in a cationic form with at least one anionic counterion. Examples of anionic counterions useful in the pharmaceutically-acceptable salts of the present invention include a halide, a carboxylate, a sulfonate, a sulfate, a phosphate, a phosphonate, a resin-bound anion, or a nitrate. When the anionic counterion is a halide, it

5 can be, for example fluoride, chloride, bromide, or iodide. Preferably the halide counterion is chloride. When the anionic counterion is a carboxylate (i.e., the anionic form of a compound containing a carboxylic acid functional group), the carboxylate counterion can vary widely. The carboxylate counterion can be, for example, formate, acetate, propionate, trifluoroacetate, succinate, salicylate, DL-aspartate, D-aspartate, L-aspartate, DL-glutamate,

10 D-glutamate, L-glutamate, glycerate, succinate, steric, DL-tartarate, D-tartarate, L-tartarate, (±)-mandelate, (R)-(-)-mandelate, (S)-(+)mandelate, citrate, mucate, maleate, malonate, benzoate, DL-malate, D-malate, L-malate, hemi-malate, 1-adamantaneacetate, 1-adamantanecarboxylate, flavianate, sulfonoacetate, (±)-lactate, L-(+)-lactate, D-(-)-lactate, pamoate, D-alpha-galacturonate, glycerate, DL-ascorbate, D-ascorbate, L-ascorbate, DL-

15 cystate, D-cystate, L-cystate, DL-homocystate, D-homocystate, L-homocystate, DL-cysteate, D-cysteate, L-cysteate, (4S)-hydroxy-L-proline, cyclopropane-1,1-dicarboxylate, 2,2-dimethylmalonate, squarate, tyrosine anion, proline anion, fumarate, 1-hydroxy-2-naphthoate, phosphonoacetate, carbonate, bicarbonate, 3-phosphonopropionate, DL-pyroglutamate, D-pyroglutamate, or L-pyroglutamate. Alternatively, the anionic counterion

20 can be a sulfonate. For example the sulfonate counterion can be methanesulfonate, toluenesulfonate, benzenesulfonate, trifluoromethylsulfonate, ethanesulfonate, (±)-camphorsulfonate, naphthalenesulfonate, 1R-(-)-camphorsulfonate, 1S-(+)-camphorsulfonate, 2-mesitylenesulfonate, 1,5-naphthalenedisulfonate, 1,2-ethanedisulfonate, 1,3-propanedisulfonate, 3-(N-morpholino)propane sulfonate, biphenylsulfonate, isethionate, or 1-hydroxy-2-naphthalenesulfonate. In another embodiment the anionic counterion can be a sulfate. Examples of sulfates useful in the present invention include without limitation sulfate, monopotassium sulfate, monosodium sulfate, and hydrogen sulfate. The anionic counterion can be a sulfamate. When the anionic counterion is a phosphate, it can be, for example, phosphate, dihydrogen phosphate, potassium hydrogen phosphate, dipotassium phosphate, potassium phosphate, sodium hydrogen phosphate, disodium phosphate, sodium phosphate, calcium dihydrogen phosphate, calcium phosphate, calcium hydrogen phosphate,

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calcium phosphate tribasic, or hexafluorophosphate. The anionic counterion can be a phosphonate. For example, the phosphonate counterion can be vinylphosphonate, 2-carboxyethylphosphonate or phenylphosphonate. Alternatively, the anionic counterion can be nitrate. The salt can also result from the addition of the compound with an oxide such as

5 zinc oxide.

The anionic counterion can, if desired, be bound to a polymeric resin. In other words, the anionic counterion can be a resin-bound anion. For example the resin-bound anion can be a polyacrylate resin wherein the resin contains anionic carboxylate groups. An example of a polyacrylate resin useful in the salts of the present invention is Bio-Rex 70

10 (produced by Bio-Rad). In an alternative example, the resin-bound anion can be a sulfonated poly (styrene divinylbenzene) copolymer resin. Non-limiting examples of sulfonated poly (styrene divinylbenzene) copolymer resins useful as anionic counterions in the present invention include Amberlite IPR-69 (Rohm & Haas) or Dowex 50WX4-400 (Dow). The polyacrylate resin or the sulfonated poly(styrene divinylbenzene) resin can be, if desired, 15 crosslinked with a crosslinking agent such as divinylbenzene.

In another embodiment, the pharmaceutically acceptable salt of the compound of Formula 1 can be one in which the present inventive compound is in an anionic form with at least one cationic counterion. The cationic counterion can be, for example, an ammonium cation, a alkali metal cation, an alkaline earth metal cation, a transition metal cation, or a 20 resin-bound cation. When the cationic counterion is an ammonium cation, it can be substituted or unsubstituted. For example, the ammonium cation can be an alkylammonium cation or a di-, tri-, or tetra-alkylammonium cation. Alternatively the ammonium cation can be an arylammonium or a di-, tri-, or tetra-aryl ammonium cation. The ammonium cation can contain both alkyl and aryl groups. The ammonium cation can be aromatic, for example 25 a pyridinium cation. Other functional groups can also be present in the ammonium cation. The ammonium cation can be, for example, ammonium, methyl ammonium, dimethylammonium, trimethylammonium, tetramethylammonium, ethanolammonium, dicyclohexylammonium, guanidinium, or ethylenediammonium cation. Alternatively the cationic counterion can be an alkali metal cation such as lithium cation, sodium cation, 30 potassium cation or cesium cation. In another alternative the cationic counterion can be an

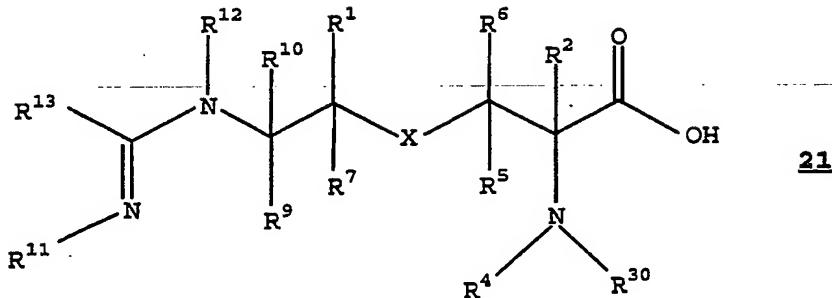
alkaline earth metal cation such as beryllium cation, magnesium cation, or calcium cation. The cation, if preferred, can be a transition metal cation such as zinc cation.

The cationic counterion can, if desired, be bound to a polymeric resin. In other words, the anionic counterion can be a resin-bound cation. For example, the resin-bound cation can be a cationically functionalized poly(styrene divinylbenzene) resin. An example of a cationically functionalized poly(styrene divinylbenzene) resin useful in the present invention is Bio-Rex-5 (Bio-Rad), an ammonium-functionalized resin. In another alternative, the resin-bound cation can be a cationically functionalized polyacrylic resin such as an aminated polyacrylic resin. An example of an aminated polyacrylic resin useful as the cationic counterion of the present invention is AG-4-XR (Bio-Rad).

In yet another embodiment of the present invention the compound of Formula 1 can be present in a zwitterionic form. In other words, the compound can contain both cationic and anionic sites within the molecule. Such a zwitterionic form can exist without a separate counterion or it can exist with both a cationic counterion and an anionic counterion.

Another embodiment provides a method for the treatment or prevention of an inflammation-related disorder wherein the method comprises treating a subject in need thereof with an inflammation-related disorder-treating or preventing amount of a compound or salt of the present invention.

In yet another embodiment the present invention provides a method for the preparation of a compound of Formula 21:



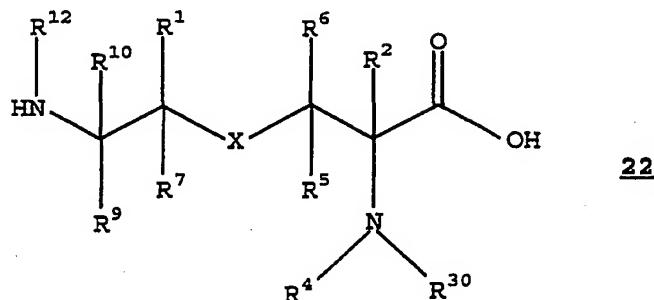
or a salt thereof, wherein: X is selected from the group consisting of -S-, -S(O)-, and -S(O)₂- ; R² is selected from the group consisting of C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-

C_5 alkoxy- C_1 alkyl, and C_1 - C_5 alkylthio- C_1 alkyl; R^{30} is selected from the group consisting of -H, -OH, -C(O)-R¹⁷, -C(O)-O-R¹⁸, and -C(O)-S-R¹⁹; R^1 , R^5 , R^6 , and R^7 independently are selected from the group consisting of -H, halogen, C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, and C_1 - C_5 alkoxy- C_1 alkyl; R^9 and R^{10} independently are selected from the group consisting of -H, C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, and C_1 - C_5 alkoxy- C_1 alkyl; with respect to R^{11} and R^{12} :

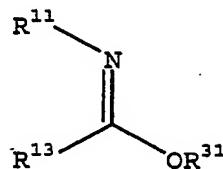
5 R^{11} is selected from the group consisting of -H, -OH, -C(O)-O-R²⁴, and -C(O)-S-R²⁵; and R^{12} is selected from the group consisting of -H, -OH, -C(O)-O-R²⁶, and -C(O)-S-R²⁷; or R^{11} is -O-, and R^{12} is -C(O)-, wherein R^{11} and R^{12} together with the atoms to which they are 10 attached form a ring; or R^{11} is -C(O)-, and R^{12} is -O-, wherein R^{11} and R^{12} together with the atoms to which they are attached form a ring;

R^{13} is C_1 alkyl;

15 R^{17} , R^{18} , R^{19} , R^{24} , R^{25} , R^{26} , R^{27} , and R^{27a} independently are selected from the group consisting of -H and alkyl, which is optionally substituted by one or more moieties selected from the group consisting of cycloalkyl, heterocyclil, aryl, and heteroaryl; and when any of R^1 , R^2 , R^4 , R^5 , R^6 , R^7 , R^9 , R^{10} , R^{11} , R^{12} , R^{13} , R^{14} , R^{15} , R^{16} , R^{17} , R^{18} , R^{19} , R^{24} , R^{25} , R^{26} , R^{27} , and R^{27a} independently is a moiety selected from the group consisting of alkyl, alkenyl, alkynyl, alkoxy, alkylthio, cycloalkyl, heterocyclil, aryl, and heteroaryl, then the moiety is optionally substituted by one or more substituent selected from the group consisting of -OH, alkoxy, and halogen. The present method of preparing compound 21 comprises treating a diamine compound having a structure corresponding to Formula 22:



(or a salt thereof) with an alkyl acetimidate having a structure corresponding to Formula 23:



23

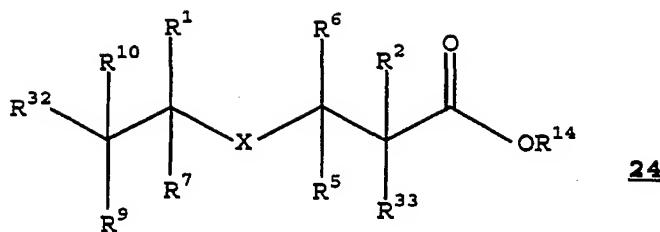
5 (or a salt thereof) wherein R^{31} is $\text{C}_1\text{-C}_6$ alkyl. Preferably R^{11} is selected from the group consisting of -H and -OH. R^{11} can, for example be -H. Alternatively, R^{11} can be -OH. When R^{11} is -H or -OH, preferably R^{13} is methyl or halomethyl. More preferably R^{13} is methyl. Also when R^{11} is -H or -OH, preferably R^{31} is $\text{C}_1\text{-C}_3$ alkyl, and more preferably ethyl. In a preferred embodiment, the treating of the diamine compound with the alkyl acetimidate is

10 performed in the presence of a base. For example, the base can be a hydrazine, a metal sulfide, a metal hydroxide, a metal alkoxide, an amine, a hydroxylamine, a metal hydride, a metal amide complex, or a basic resin. When the base is a basic resin it can be, for example a polymer-bound diazabicyclo[4.4.0]dec-2-ene resin. For example, the basic resin can have a poly(styrene divinylbenzene) copolymer backbone with diazabicyclo[4.4.0]dec-2-ene

15 bonded to the copolymer. When the base is an amine, it can be essentially any substituted or unsubstituted amine. For example, the amine can be 1,5-diazabicyclo[4.3.0]non-5-ene; 1,4-diazabicyclo[2.2.2]octane; or 1,8-diazabicyclo[5.4.0]undec-7-ene. When the base is an alkali metal hydroxide it can be, for example, potassium hydroxide or sodium hydroxide. When the base is a metal hydride it can be, for example, sodium hydride, potassium hydride, or

20 calcium hydride.

In a further embodiment of the present invention provides a method for the preparation of a diamine compound having a structure corresponding to Formula 22 (or a salt thereof), wherein the method comprises treating a protected diamine compound having the structure corresponding to Formula 24:



(or a salt thereof) wherein R^{33} is selected from the group consisting of $-NH_2$ and a protected amino group; and R^{32} is a protected amino group; and R^{14} is selected from the group

5 consisting of $-H$ and C_1-C_6 alkyl; wherein when R^{14} is C_1-C_6 alkyl, R^{14} is optionally substituted by one or more moieties selected from the group consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl; wherein the treating is performed with a deprotecting reagent, thereby producing the diamine compound. Protected amino groups useful in the present invention vary widely in nature. Numerous protected amino groups useful in the

10 present invention for either R^{32} or R^{33} are described by Theodora W. Greene and Peter G.M. Wuts (Protective Groups in Organic Synthesis, 3rd ed., John Wiley & Sons, New York, 1999, pp. 494-653). For example either or both of R^{32} and R^{33} can be a 4-chlorobenzylimino group. When R^{33} is a 4-chlorobenzylimino group, preferably R^{32} is $-NH_2$. In another embodiment, either or both of R^{32} and R^{33} can be a t-butoxycarbonylamino group. When R^{33}

15 is a t-butoxycarbonylamino group, preferably R^{32} is $-NH_2$. In yet another embodiment, either or both of R^{32} and R^{33} can be an N-phthalimido group. When R^{33} is an N-phthalimido group, preferably R^{32} is $-NH_2$. In another embodiment, either or both of R^{32} and R^{33} can be a benzyloxycarbonylamino group. In one preferred embodiment of the present invention, the protected amino group is any such group resulting from the reaction of an aldehyde with the

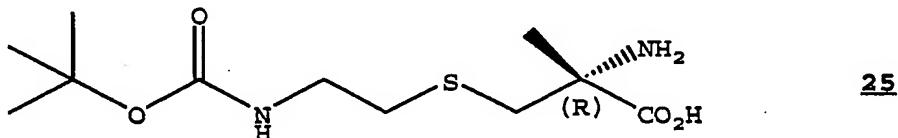
20 corresponding amino group to form a Schiff base. A large variety of deprotecting reagents can be advantageously used in the present invention to effect the conversion of 24 to 22. Many such deprotecting reagents are described by Greene and Wuts, *supra*. For example, when the protected amino group is a 4-chlorobenzylimino group or a t-butoxycarbonylamino group, preferably the deprotecting reagent is an acid. Some useful acid deprotecting agents

25 include without limitation hydrochloric acid, hydrobromic acid, sulfuric acid, trifluoroacetic acid, phosphoric acid, phosphorus acid, and acetic acid. In another example, when R^{32} or R^{33} is an N-phthalimido group, the deprotecting reagent can be either an acid or a base. When

the deprotecting reagent for the N-phthalimido group is a base, the base can be, for example, a hydrazine, a metal sulfide, a metal hydroxide, a metal alkoxide, an amine, a hydroxylamine, and a metal amide complex. When the deprotecting reagent for the N-phthalimido group is an acid, the acid can be, for example, hydrochloric acid, hydrobromic acid, hydroiodic acid, sulfuric acid, methanesulfonic acid, trifluoromethanesulfonic acid, toluenesulfonic acid, benzenesulfonic acid, trifluoroacetic acid, phosphoric acid, phosphorus acid, or acetic acid. Preferably the treating of compound 24 with the deprotecting reagent is performed in the presence of water.

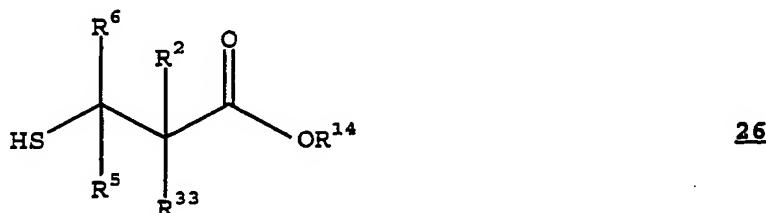
In the present reaction, R^{14} is preferably -H. R^{33} preferably is -NH₂, or a salt thereof. 10 R^2 is preferably methyl. In another preferred embodiment R^1 , R^5 , R^6 , R^7 , R^9 , and R^{10} each is -H. A further preferred embodiment is one in which R^{33} is a t-butoxycarbonylamino group. In a particularly preferred embodiment, compound 24 has the structure corresponding to Formula 25:

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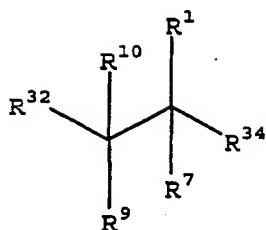


(or a salt thereof) wherein the parenthetical R means that the carbon alpha to the carboxylic acid function is in the R absolute configuration. In other words, compound 25 is the R-enantiomer or a salt thereof.

20 The present invention also provides a method for the preparation of the protected diamine compound 24 (or a salt thereof) wherein the method comprises treating a sulphydryl compound having the structure of Formula 26:



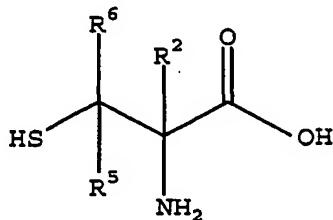
25 with a protected amino ethyl alkylating compound having the structure of Formula 27:



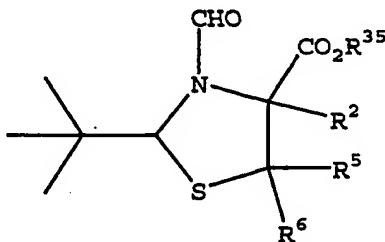
wherein R³⁴ is a nucleophilic substitution leaving group; thereby forming the protected diamine compound. Preferably, this reaction is performed in the presence of a base. The base can be, for example, a hydrazine, a metal sulfide, a metal hydroxide, a metal alkoxide, 5 an amine, a hydroxylamine, and a metal amide complex. Preferably the base is an alkali metal hydroxide and more preferably the base is potassium hydroxide or sodium hydroxide. R³⁴ in the structure of Formula 27 can vary widely and can represent essentially any nucleophilic leaving group which produces either a pharmaceutically acceptable anion or an anion which can be exchanged for a pharmaceutically acceptable anion. In other words,

10 (R³⁴)⁻ is a pharmaceutically acceptable anion or an anion which can be exchanged for a pharmaceutically acceptable anion. For example, R³⁴ can be chloro, bromo, iodo, methanesulfonato, toluenesulfonato, benzenesulfonato, or trifluoromethanesulfonato. Preferably R³⁴ is chloro, bromo, or iodo and more preferably R³⁴ is bromo. In the present reaction it is preferred for R³³ to be -NH₂. In the present reaction R² can be C₁-C₆ alkyl 15 optionally substituted by one or more substituent selected from the group consisting of -OH, alkoxy and halogen. Preferably R² is C₁-C₃ alkyl optionally substituted by one or more substituent selected from the group consisting of -OH, alkoxy and halogen, and more preferably R² is C₁-C₃ alkyl. For example, R² advantageously can be methyl. In a further embodiment R⁵ and R⁶ each can be -H. In yet another embodiment R¹ and R⁷ independently 20 can be selected from the group consisting of H and C₁-C₆ alkyl optionally substituted by one or more halogen; preferably R¹ and R⁷ each is -H. In the present reaction, R³² preferably can be selected from the group consisting of a 4-chlorobenzylimino group, a t-butoxycarbonylamino group, and an N-phthalimido group, and more preferably R³² is a t-butoxycarbonylamino group.

The present invention also provides a method for the preparation of a sulphydryl compound having the structure of Formula 28:

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wherein R², R⁵, and R⁶ are as defined above, and wherein the method comprises treating 5 under hydrolysis conditions a thiazolidine compound having the structure of Formula 29:

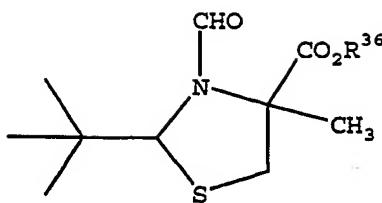
29

or a salt thereof, wherein R³⁵ is a moiety selected from the group consisting of -H and C₁-C₆ alkyl; thereby forming the sulphydryl compound. The hydrolysis conditions preferably comprise contacting the thiazolidine compound with an acid in the presence of water.

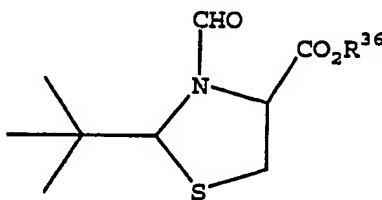
10 Preferably the acid is selected from the group consisting of hydrochloric acid, hydrobromic acid, hydroiodic acid, sulfuric acid, methanesulfonic acid, trifluoromethanesulfonic acid, toluenesulfonic acid, benzenesulfonic acid, trifluoroacetic acid, phosphoric acid, phosphorus acid, and acetic acid. In one embodiment of the present reaction R⁵ and R⁶ each is -H. In another embodiment R² of compound 29 is C₁-C₆ alkyl optionally substituted by one or

15 more substituent selected from the group consisting of -OH, alkoxy, and halogen. Preferably R² is C₁-C₃ alkyl optionally substituted by one or more substituent selected from the group consisting of alkoxy and halogen, and more preferably R² is C₁-C₃ alkyl. For example, R² can be methyl. In another embodiment of the present invention R³⁵ of compound 29 is methyl.

20 The present invention also provides a method for the preparation of a methyl thiazolidine compound having the structure of Formula 30:

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(or a pharmaceutically-acceptable salt thereof) wherein R^{36} is C_1 - C_6 alkyl; wherein the method comprises treating under methylation conditions a deprotonatable thiazolidine compound having the structure of Formula 31:

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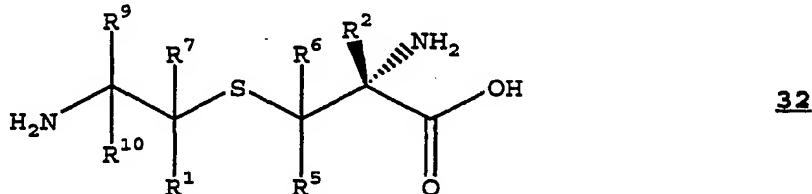
thereby forming the methyl thiazolidine compound. Preferably the methylation conditions comprise treating the deprotonatable thiazolidine compound with a base and a methylating agent. The nature of the base can vary widely. The base can be, for example, a metal hydroxide, a metal alkoxide, a metal hydride, a metal alkyl, and a metal amide complex.

10 Preferably the base is a metal amide complex. Some metal amide complexes useful as a base in the present invention include without limitation lithium hexamethyldisilazide, sodium hexamethyldisilazide, potassium hexamethyldisilazide, lithium diisopropyl amide, sodium diisopropyl amide, potassium diisopropyl amide, sodium amide, lithium amide, potassium amide, sodium diethylamide, lithium diethylamide, potassium diethylamide, methyl lithium, 15 t-butyl lithium, sec-butyl lithium, methyl sodium, t-butyl sodium, sec-butyl sodium, and methyl magnesium bromide. In the present methylation reaction R^{36} can be C_1 - C_3 alkyl; for example R^{36} can be methyl.

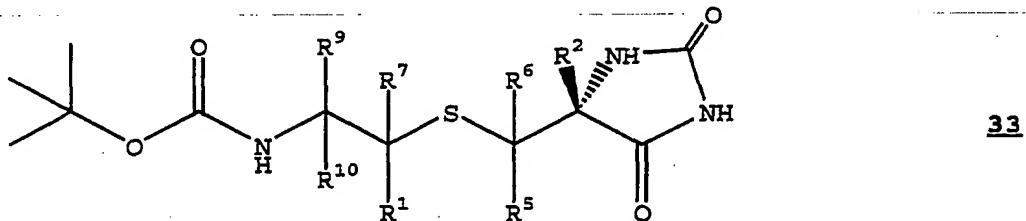
20 The present invention further provides a method for the preparation of the deprotonatable thiazolidine compound 31 wherein the method comprises contacting under condensing conditions a cysteine C_1 - C_6 alkyl ester with pivalaldehyde, thereby forming the deprotonatable thiazolidine compound. Preferably condensing conditions comprise performing the contacting in the presence of a base. The base can vary widely. For example the base can be, without limitation, a hydrazine, a metal sulfide, a metal hydroxide, a metal

alkyl base, a metal alkoxide, an amine, a hydroxylamine, and a metal amide complex. When the base is a metal amide complex, it can be, for example, lithium bis(trimethylsilyl)amide. In the present condensation reaction it is preferred that R^{36} is C_1 - C_3 alkyl, and more preferably R^{36} is methyl.

5 In another embodiment the present invention provides a method for the preparation of an alpha-amino acid compound having the structure of Formula 32:

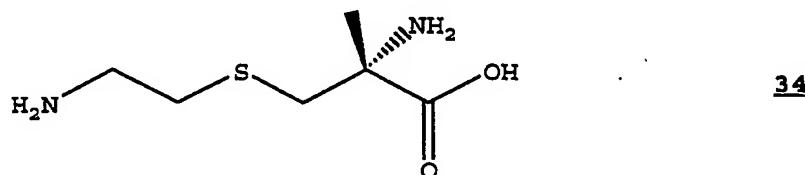


(or a salt, an enantiomer, or a racemate thereof) wherein: R^2 is selected from the group consisting of C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_1 - C_5 alkoxy- C_1 alkyl, and C_1 - C_5 alkylthio- C_1 alkyl; R^1 , R^5 , R^6 , and R^7 independently are selected from the group consisting of -H, halogen, C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, and C_1 - C_5 alkoxy- C_1 alkyl; R^9 and R^{10} independently are selected from the group consisting of -H, C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, and C_1 - C_5 alkoxy- C_1 alkyl; and when any of R^1 , R^2 , R^5 , R^6 , R^7 , R^9 , and R^{10} independently is a moiety selected from the group consisting of alkyl, alkenyl, and alkynyl, then the moiety is optionally substituted by one or more substituent selected from the group consisting of -OH, alkoxy, and halogen; wherein the method comprises treating under hydrolyzing conditions a hydantoin compound having the structure of Formula 33:



(or a salt, an enantiomer, or a racemate thereof), thereby forming the alpha-amino acid compound. The hydrolyzing conditions can comprise, for example, contacting the hydantoin compound with an acid to produce an acid hydrolyzate. Acids useful in the present hydrolysis reaction include, for example, hydrochloric acid, hydrobromic acid, hydroiodic

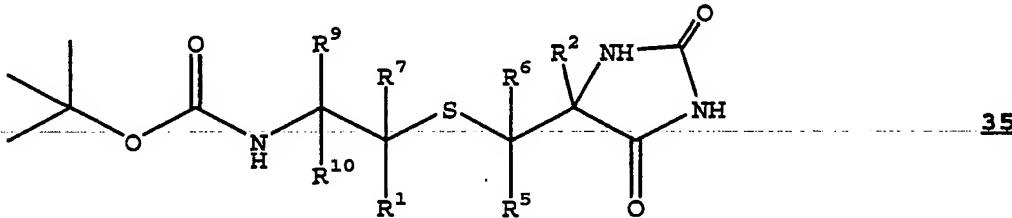
acid, sulfuric acid, trifluoroacetic acid, or phosphoric acid. The method for the preparation of alpha-amino acid compound 32 can further comprise treating the acid hydrolyzate with an ion exchange resin. Alternatively, the hydrolyzing conditions can comprise contacting the hydantoin compound with a base to produce a base hydrolyzate. Bases useful in the present 5 base hydrolysis include without limitation a hydrazine, a metal sulfide, a metal hydroxide, or a metal alkoxide. Whether the hydrolysis is base-mediated or acid-mediated, it is preferred that R¹, R⁵, R⁶, and R⁷ each is -H in the structure of compound 33. It is also preferred that R⁹ and R¹⁰ each is -H. In a particularly preferred embodiment, the alpha-amino acid compound has the structure of Formula 34:



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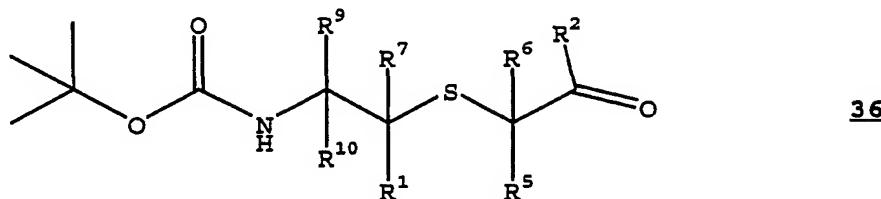
(or a salt, an enantiomer, or a racemate thereof).

The present invention further provides a method for the preparation of a hydantoin compound having the structure of Formula 35:



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(or a salt, an enantiomer, or a racemate thereof) wherein R¹, R², R⁵, R⁶, R⁷, R⁹, and R¹⁰ are as defined above, wherein the method comprises contacting an alpha-sulfo ketone compound having the structure of Formula 36:

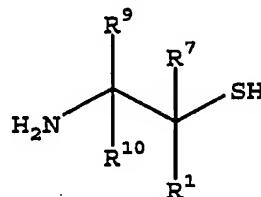


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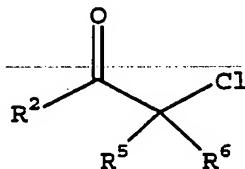
with a source of cyanide in the presence of a source of ammonium carbonate and water, thereby producing the hydantoin compound. The source of cyanide can be, for example, hydrogen cyanide or a metal cyanide salt. When the source of cyanide is a metal cyanide

salt, preferably the salt is sodium cyanide, potassium cyanide, or lithium cyanide. More preferably the metal cyanide salt is sodium cyanide. For compound 36 in the present hydantoin-forming reaction, R¹, R⁵, R⁶, and R⁷ each is preferably -H. It is further preferred that R⁹ and R¹⁰ each is -H. The method of preparing compound 35 can further comprise a 5 chiral separation step. When the method of preparing compound 35 further comprises a chiral separation step, then the hydantoin compound product preferably has the structure of compound 33 (or a salt or an enantiomer thereof).

10 Also provided by the present invention is a method for the preparation of an alpha-sulfo ketone compound 36 wherein the method comprises contacting an aminothiol compound having the structure of Formula 37:

37

15 with di-t-butyl carbonate in the presence of a base to produce an intermediate mixture; and contacting the intermediate mixture with an alpha-chloro ketone compound having the structure of Formula 38:

38

thereby producing the alpha-sulfo ketone compound. In the present reaction the base can vary widely. For example, the base can be without limitation a hydrazine, a metal sulfide, a metal hydroxide, a metal alkoxide, an amine, a hydroxylamine, and a metal amide complex. Preferably the base is a metal hydroxide such as sodium hydroxide, potassium hydroxide, or 20 lithium hydroxide. In the present reaction it is preferred that R¹, R⁵, R⁶, and R⁷ each is -H. It is further preferred that R⁹ and R¹⁰ each is -H.

25 The term "pharmaceutically-acceptable salts" embraces salts commonly used to form alkali metal salts and to form addition salts of free acids or free bases. The nature of the salt is not critical, provided that it is pharmaceutically acceptable. Pharmaceutically acceptable salts are particularly useful as products of the methods of the present invention because of

their greater aqueous solubility relative to a corresponding parent or neutral compound. Such salts must have a pharmaceutically acceptable anion or cation. Suitable pharmaceutically-acceptable acid addition salts of compounds of the present invention may be prepared from inorganic acid or from an organic acid. Examples of such inorganic acids 5 are hydrochloric, hydrobromic, hydroiodic, nitric, carbonic, sulfuric and phosphoric acid. Appropriate organic acids include from aliphatic, cycloaliphatic, aromatic, araliphatic, heterocyclic, carboxylic and sulfonic classes of organic acids, examples of which are formic, acetic, propionic, succinic, glycolic, gluconic, lactic, malic, tartaric, citric, ascorbic, glucoronic, maleic, fumaric, pyruvic, aspartic, glutamic, benzoic, anthranilic, mesylic, 10 salicylic, p-hydroxybenzoic, phenylacetic, mandelic, embonic (parnoic), methanesulfonic, ethylsulfonic, benzenesulfonic, sulfanilic, stearic, cyclohexylaminosulfonic, algenic, galacturonic acid. Suitable pharmaceutically-acceptable base addition salts of compounds of the present invention include metallic salts made from aluminum, calcium, lithium, magnesium, potassium, sodium and zinc or organic salts made from N,N'- 15 dibenzylethylenediamine, choline, chlorprocaine, diethanolamine, ethylenediamine, meglumine (N-methylglucamine) and procain. Suitable pharmaceutically acceptable acid addition salts of the compounds of the present invention when possible include those derived from inorganic acids, such as hydrochloric, hydrobromic, boric, fluoroboric, phosphoric, metaphosphoric, nitric, carbonic (including carbonate and hydrogen carbonate anions), 20 sulfonic, and sulfuric acids, and organic acids such as acetic, benzenesulfonic, benzoic, citric, ethanesulfonic, fumaric, gluconic, glycolic, isothionic, lactic, lactobionic, maleic, malic, methanesulfonic, trifluoromethanesulfonic, succinic, toluenesulfonic, tartaric, and trifluoroacetic acids. Suitable pharmaceutically acceptable base salts include ammonium salts, alkali metal salts such as sodium and potassium salts, and alkaline earth salts such as 25 magnesium and calcium salts. All of these salts may be prepared by conventional means from the corresponding conjugate base or conjugate acid of the compounds of the present invention by reacting, respectively, the appropriate acid or base with the conjugate base or conjugate acid of the compound. Another pharmaceutically acceptable salt is a resin-bound salt.

30 While it may be possible for the compounds of the present invention to be administered as the raw chemical, it is preferable to present them as a pharmaceutical

composition. According to a further aspect, the present invention provides a pharmaceutical composition comprising a compound of the present invention or a pharmaceutically acceptable salt or solvate thereof, together with one or more pharmaceutically acceptable carriers thereof and optionally one or more other therapeutic ingredients. The carrier(s) must

5 be acceptable in the sense of being compatible with the other ingredients of the formulation and not deleterious to the recipient thereof.

The formulations include those suitable for oral, parenteral (including subcutaneous, intradermal, intramuscular, intravenous and intraarticular), rectal and topical (including dermal, buccal, sublingual and intraocular) administration although the most suitable route

10 may depend upon for example the condition and disorder of the recipient. The formulations may conveniently be presented in unit dosage form and may be prepared by any of the methods well known in the art of pharmacy. All methods include the step of bringing into association a compound of The present invention or a pharmaceutically acceptable salt or solvate thereof with the carrier which constitutes one or more accessory ingredients. In

15 general, the formulations are prepared by uniformly and intimately bringing into association the active ingredient with liquid carriers or finely divided solid carriers or both and then, if necessary, shaping the product into the desired formulation.

Formulations of the present invention suitable for oral administration may be presented as discrete units such as capsules, cachets or tablets each containing a

20 predetermined amount of the active ingredient; as a powder or granules; as a solution or a suspension in an aqueous liquid or a non-aqueous liquid; or as an oil-in-water liquid emulsion or a water-in-oil liquid emulsion. The active ingredient may also be presented as a bolus, electuary or paste.

A tablet may be made by compression or molding, optionally with one or more

25 accessory ingredients. Compressed tablets may be prepared by compressing in a suitable machine the active ingredient in a free-flowing form such as a powder or granules, optionally mixed with a binder, lubricant, inert diluent, lubricating, surface active or dispersing agent. Molded tablets may be made by molding in a suitable machine a mixture of the powdered compound moistened with an inert liquid diluent. The tablets may optionally be coated or

30 scored and may be formulated so as to provide slow or controlled release of the active ingredient therein.

Formulations for parenteral administration include aqueous and non-aqueous sterile injection solutions which may contain anti-oxidants, buffers, bacteriostats and solutes which render the formulation isotonic with the blood of the intended recipient; and aqueous and non-aqueous sterile suspensions which may include suspending agents and thickening agents. The formulations may be presented in unit-dose or multi-dose containers, for example sealed ampoules and vials, and may be stored in a freeze-dried (lyophilized) condition requiring only the addition of the sterile liquid carrier, for example, saline, water-for-injection, immediately prior to use. Extemporaneous injection solutions and suspensions may be prepared from sterile powders, granules and tablets of the kind previously described.

10 Formulations for rectal administration may be presented as a suppository with the usual carriers such as cocoa butter or polyethylene glycol.

Formulations for topical administration in the mouth, for example buccally or sublingually, include lozenges comprising the active ingredient in a flavored basis such as sucrose and acacia or tragacanth, and pastilles comprising the active ingredient in a basis

15 such as gelatin and glycerin or sucrose and acacia.

Preferred unit dosage formulations are those containing an effective dose, as hereinbelow recited, or an appropriate fraction thereof, of the active ingredient.

It should be understood that in addition to the ingredients particularly mentioned above, the formulations of this invention may include other agents conventional in the art

20 having regard to the type of formulation in question, for example those suitable for oral administration may include flavoring agents.

The compounds of the invention may be administered orally or via injection at a dose of from 0.001 to 2500 mg/kg per day. The dose range for adult humans is generally from 0.005 mg to 10 g/day. Tablets or other forms of presentation provided in discrete units may

25 conveniently contain an amount of compound of the invention which is effective at such dosage or as a multiple of the same, for instance, units containing 5 mg to 500 mg, usually around 10 mg to 200 mg.

The compounds of Formula 1 are preferably administered orally or by injection (intravenous or subcutaneous). The precise amount of compound administered to a patient

30 will be the responsibility of the attendant physician. However, the dose employed will depend on a number of factors, including the age and sex of the patient, the precise disorder

being treated, and its severity. Also, the route of administration may vary depending on the condition and its severity.

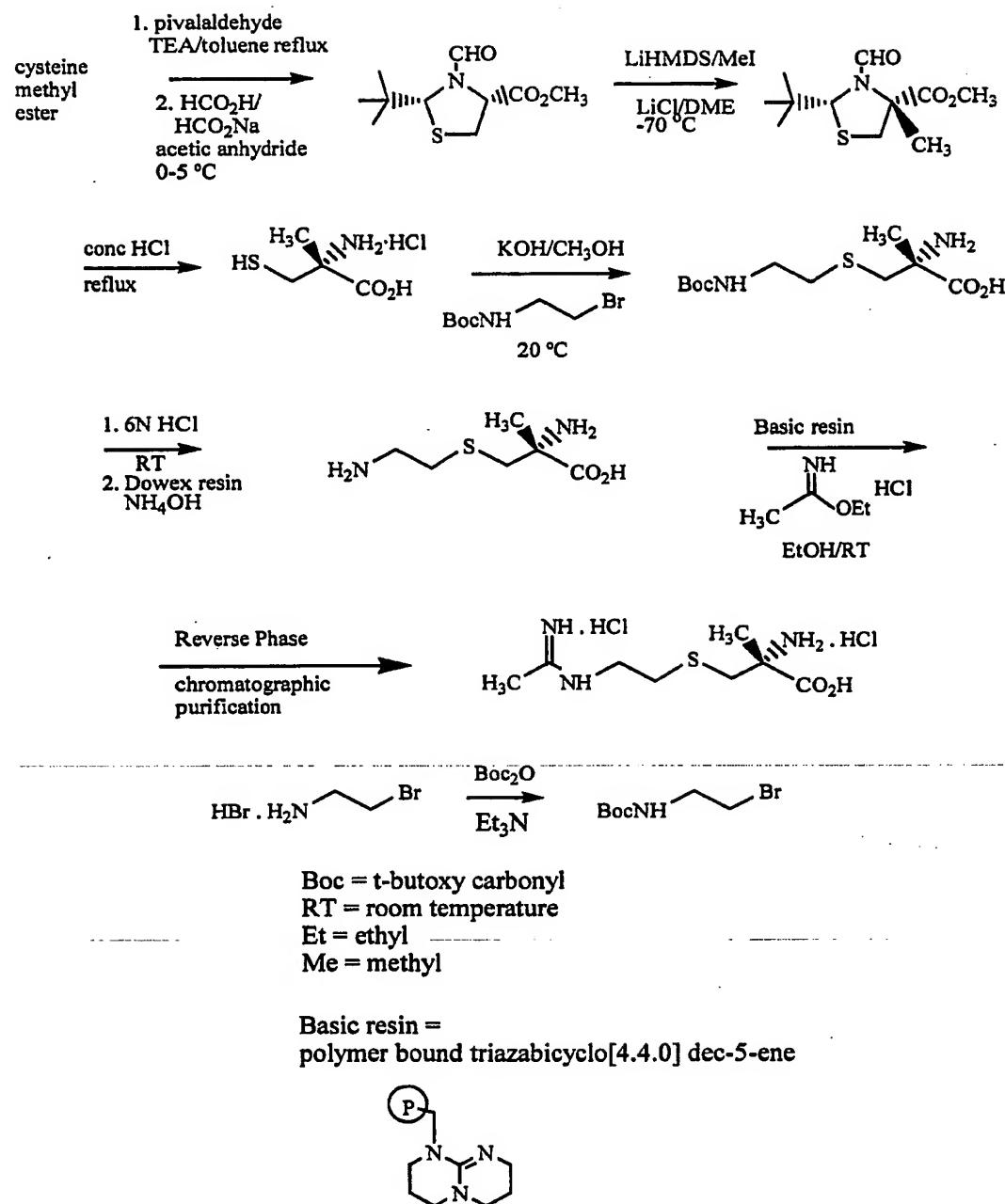
Compounds of the present invention can exist in tautomeric, geometric or stereoisomeric forms. The present invention contemplates all such compounds, including 5 cis- and trans-geometric isomers, E- and Z-geometric isomers, R- and S-enantiomers, diastereomers, d-isomers, l-isomers, the racemic mixtures thereof and other mixtures thereof, as falling within the scope of the invention. Pharmaceutically acceptable sales of such tautomeric, geometric or stereoisomeric forms are also included within the invention.

The terms "cis" and "trans" denote a form of geometric isomerism in which two 10 carbon atoms connected by a double bond will each have two highest ranking groups on the same side of the double bond ("cis") or on opposite sides of the double bond ("trans"). Some of the compounds described contain alkenyl groups, and are meant to include both cis and trans or "E" and "Z" geometric forms.

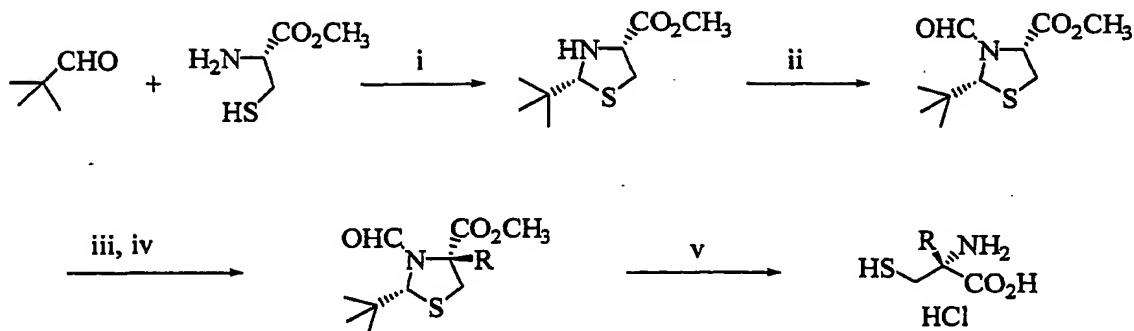
Some of the compounds described contain one or more stereocenters and are meant 15 to include R, S, and mixtures of R and S forms for each stereocenter present.

The following general synthetic sequences are useful in making the present invention.

Scheme 1a



Scheme 1b

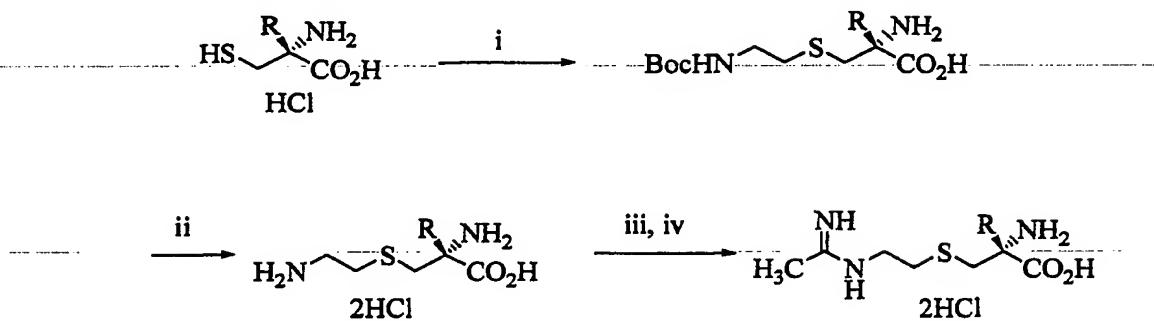


R = alkyl or alkoxyalkyl

(i) pentane with Dean-Stark trap; (ii) HCO₂Na, Ac₂O, HCO₂H;
 (iii) LiN[(CH₃)₃Si]₂, DMPU, THF, -78 °C;
 (iv) RI or R-SO₃CF₃; (v) 6N HCl, reflux 2d.

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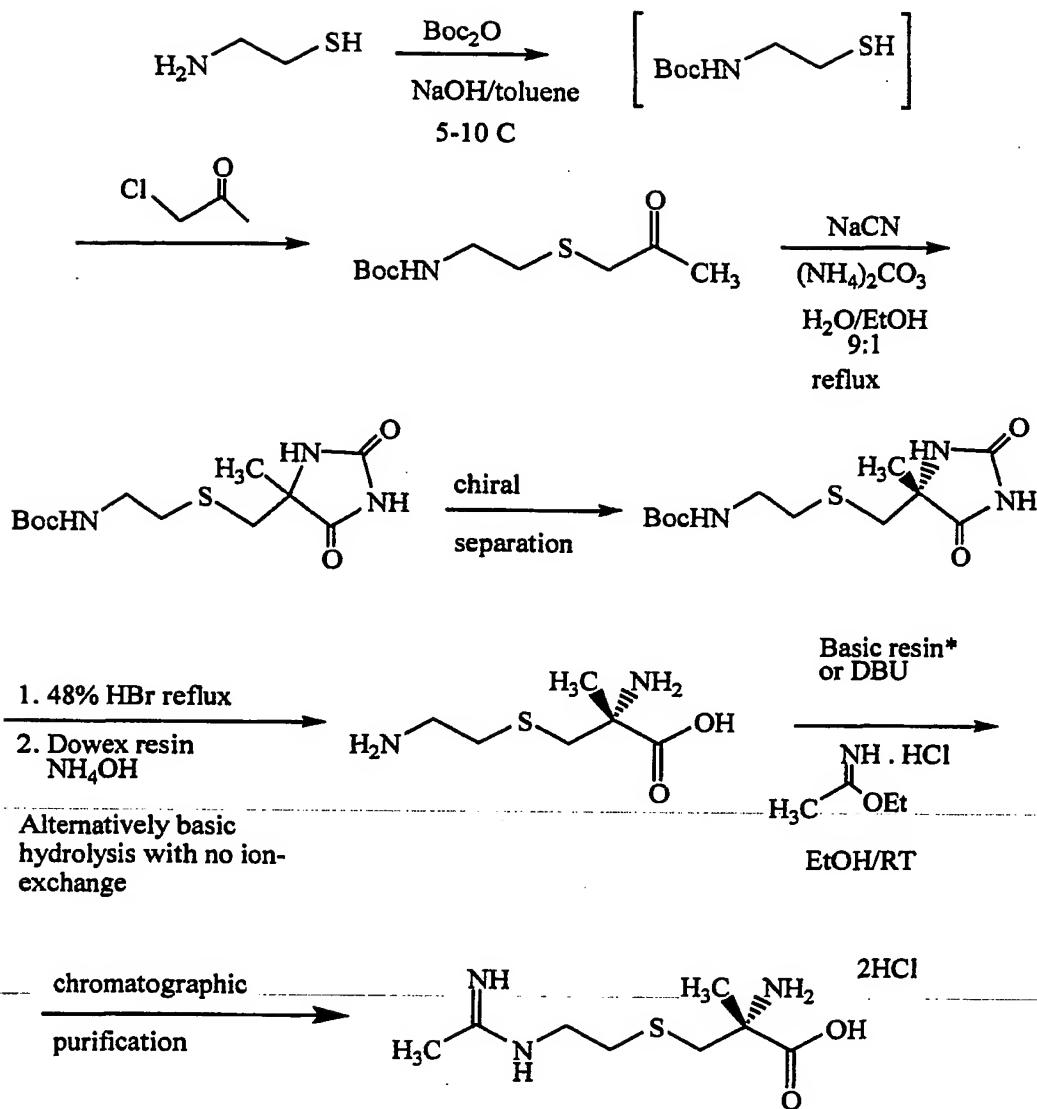
Scheme 1c

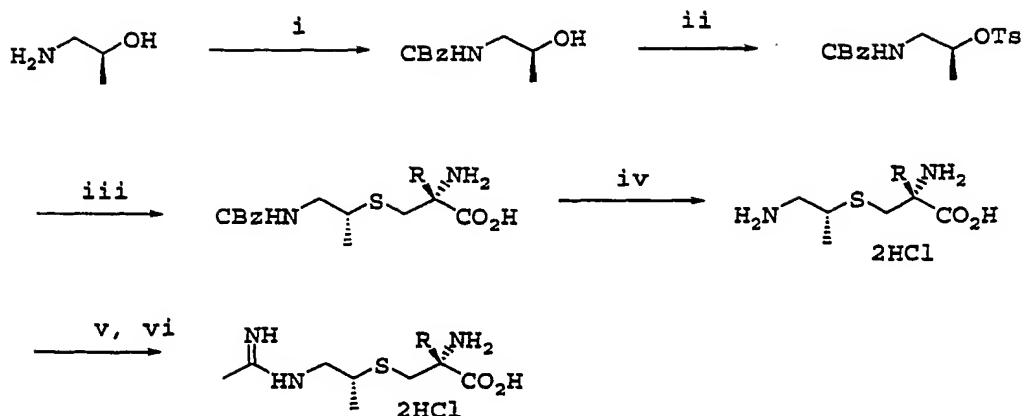


R = alkyl or alkoxyalkyl

(i) NaH, NMP, Boc-NHCH₂CH₂Br; (ii) 1NHCl;
 (iii) ethylacetimidate, NaOH; (iv) ion exchange

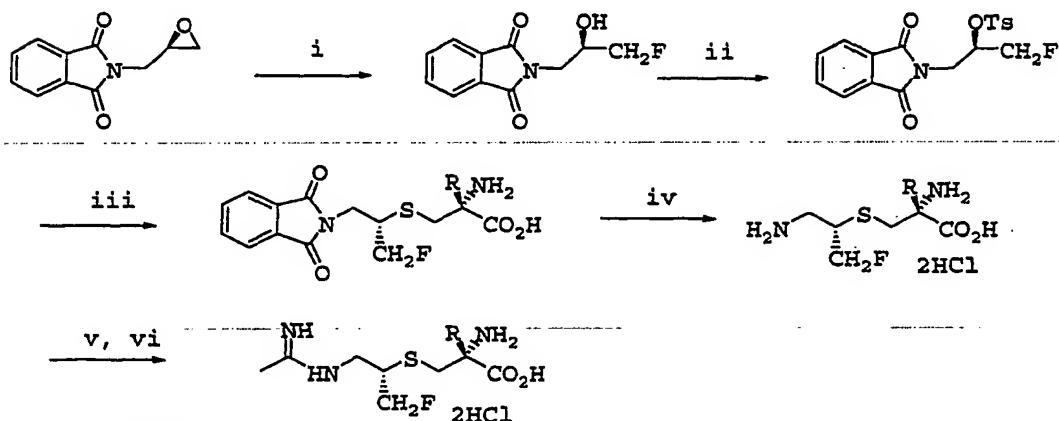
Scheme 2



Scheme 3*R* = alkyl

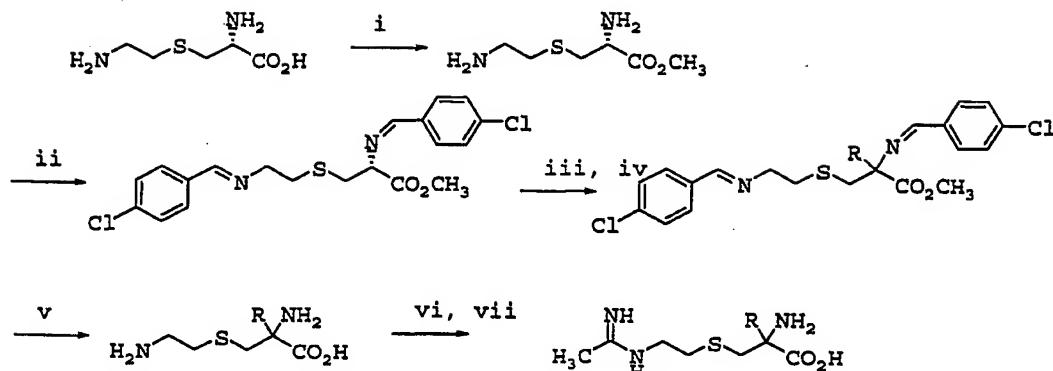
(i) Benzylchloroformate, benzene; (ii) Toluene sulfonyl chloride, triethylamine; (iii) 2-methyl-L-cysteine-HCl, NaH, NMP; (iv) 6N HCl, reflux; (v) ethylacetimidate, NaOH; (vi) ion exchange.

5

Scheme 4*R* = alkyl

(i) KHF_2 , $\text{nBu}_4\text{NH}_2\text{F}_3$; (ii) Toluene sulfonyl chloride, triethylamine; (iii) 2-methyl-L-cysteine-HCl, NaH, NMP; (iv) 6N HCl, reflux; (v) ethylacetimidate, NaOH; (vi) ion exchange.

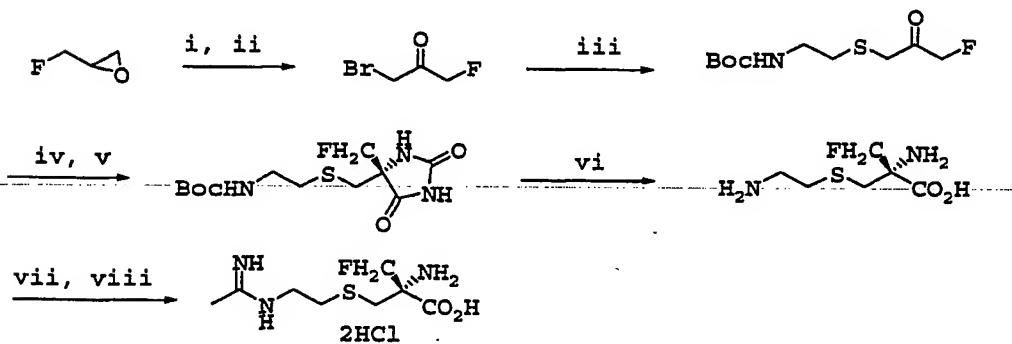
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Scheme 5

R = alkyl or alkoxyalkyl

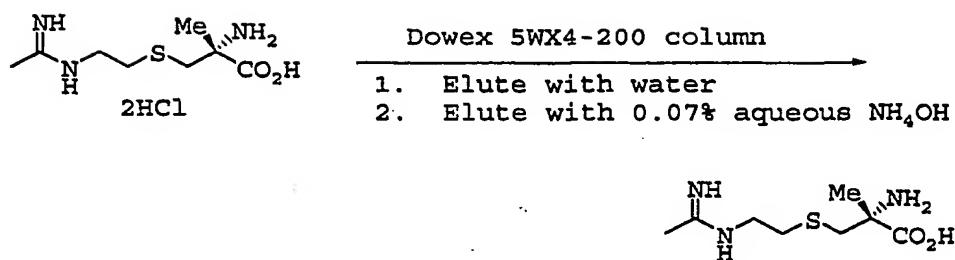
(i) CH_3OH , HCl ; (ii) 4-chlorobenzaldehyde, $(\text{CH}_3\text{CH}_2)_3\text{N}$, CH_3CN , MgSO_4 ; (iii) $\text{NaN}[(\text{CH}_3)_3\text{Si}]_2$, -78°C ; (iv) RI ; (v) HCl ; (vi) ethyl acetimidate, base; (vii) ion exchange.

5

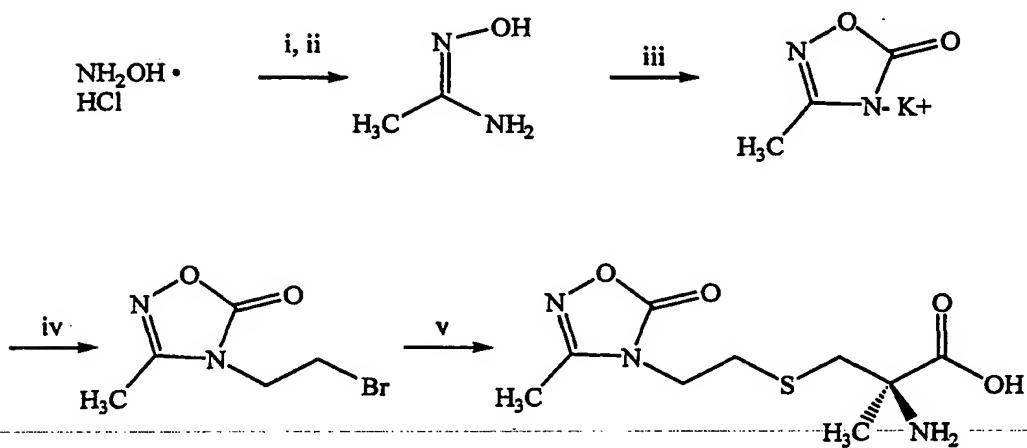
Scheme 6

(i) HBr (Synthesis 1971 646-7, J Chem Soc 1961 3448-52); (ii) $\text{K}_2\text{Cr}_2\text{O}_7$; (iii) $\text{Boc-NHCH}_2\text{CH}_2\text{SH}$, NaOH , toluene; (iv) NaCN , $(\text{NH}_4)_2\text{CO}_3$, EtOH ; (v) Chiral Separation; (vi) 48% HBr ; (vii) ethyl acetimidate, base; (viii) HCl .

10

Scheme 7

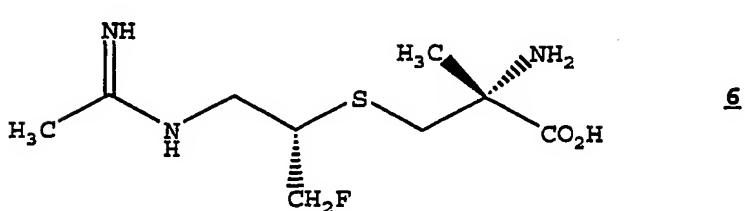
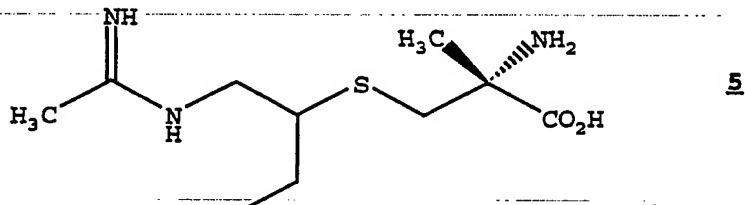
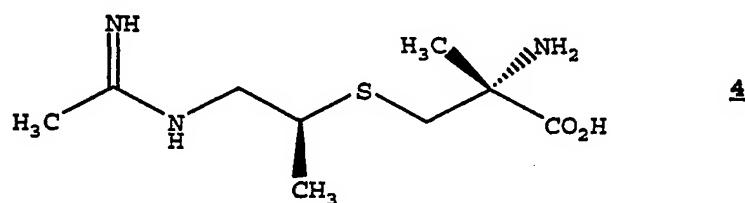
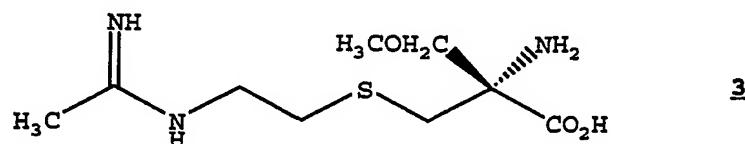
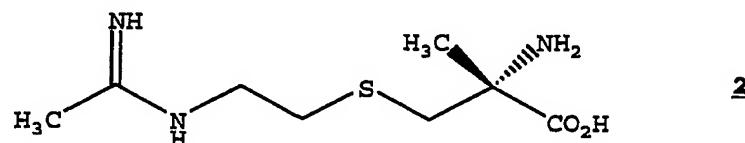
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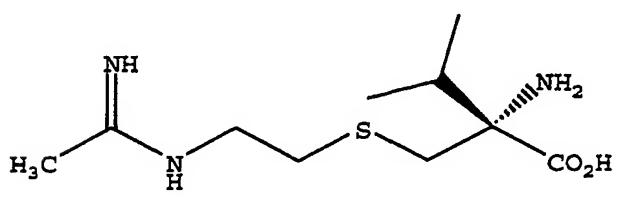
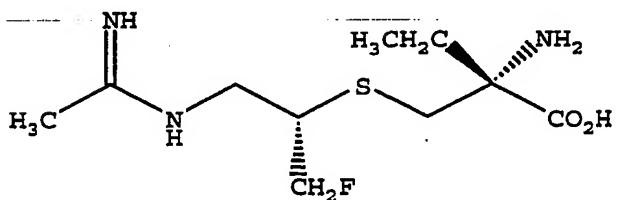
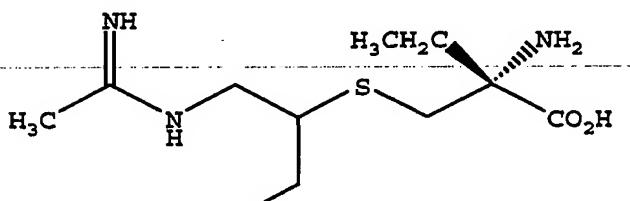
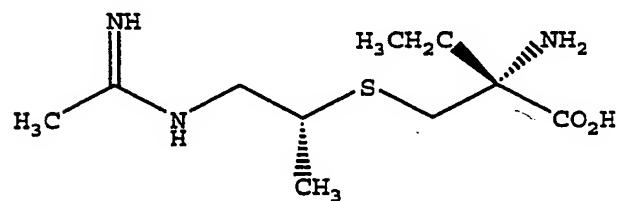
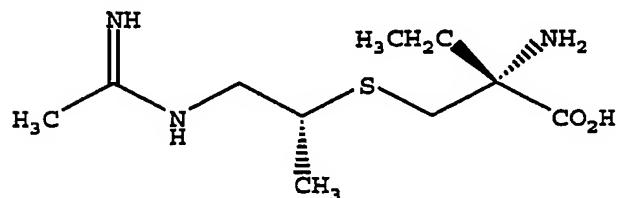
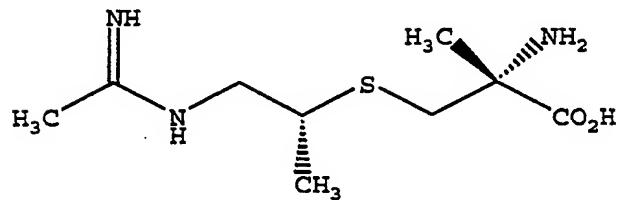
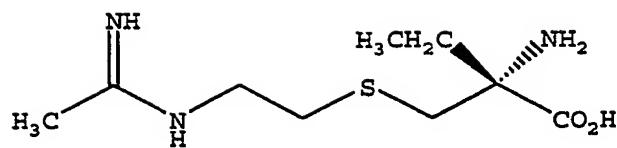
Scheme 8

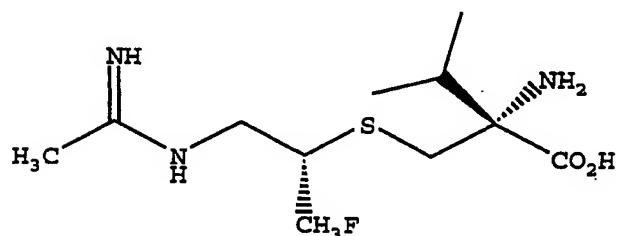
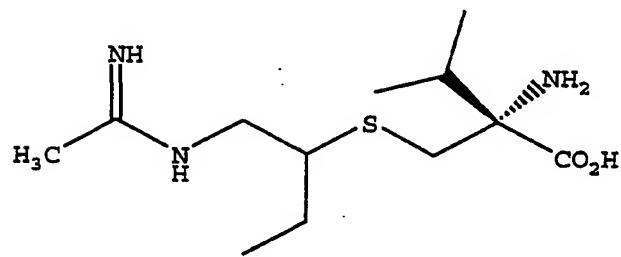
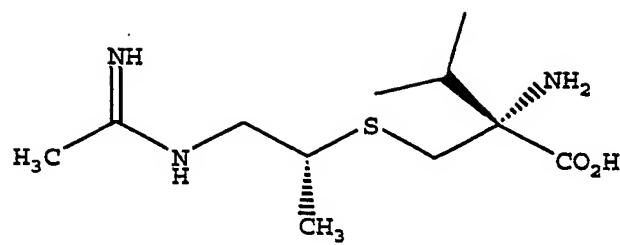
i) NaOH; ii) acetonitrile, reflux overnight; iii) diethyl carbonate, potassium t-butoxide;
 iv) 1,2-dibromoethane, DMF; v) NaOH, α -methyl cysteine

The following structures are illustrative of the many compounds provided by the present invention. The examples provided here are not intended to be limiting and one of skill in the art, given the present disclosure, will recognize that many alternative structures 5 are embraced by the present invention.

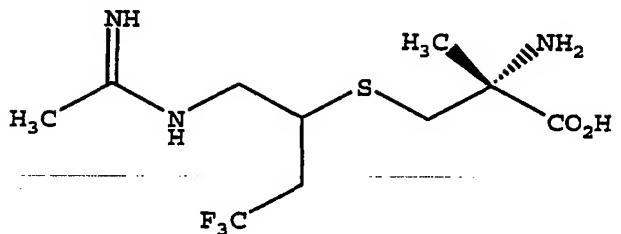
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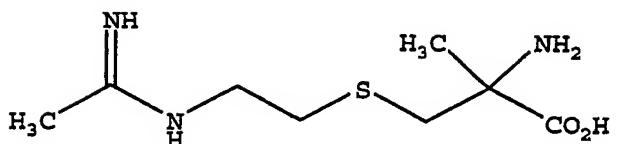


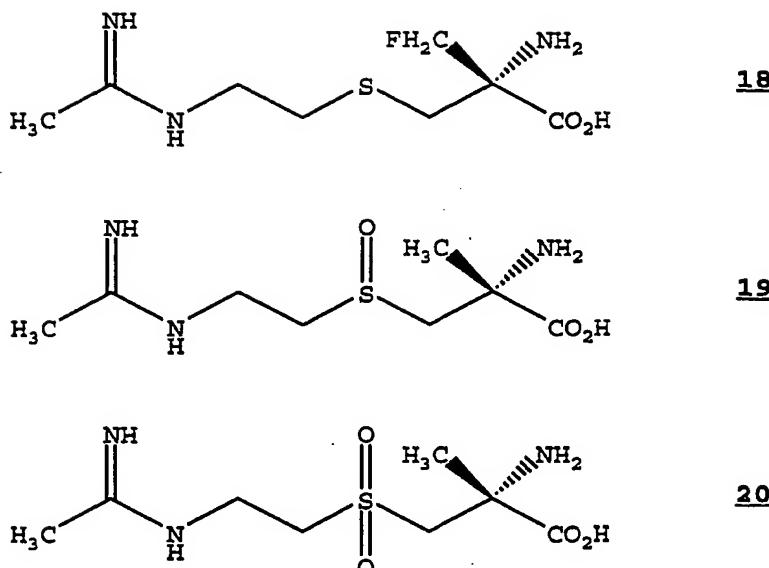


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The following compounds are further examples of compounds embraced by the present invention or useful in the preparation of compounds of the present invention.

10

(2*R*,4*R*)-Methyl-2-tert-butyl-1,3-thiazoline-3-formyl-4-carboxylate.

(2*R*,4*R*)-Methyl-2-tert-butyl-1,3-thiazoline-3-formyl-4-methyl-4-carboxylate.

S-[2-[(1,1-Dimethylethoxy)carbonyl]amino]ethyl]-2-methyl-L-cysteine trifluoroacetate.

15 (S)-1-[(Benzylloxycarbonyl)amino]-2-propanol.

(S)-1-[(Benzylloxycarbonyl)amino]-2-propanol tosylate.

S-[(1*R*)-2-(Benzylloxycarbonylamino)-1-methylethyl]-2-methyl-L-cysteine trifluoroacetate.

S-[(1*R*)-2-Amino-1-methylethyl]-2-methyl-L-cysteine hydrochloride.

20 *S*-(2-Aminoethyl)-L-cysteine, methyl ester.

N-{4-Chlorophenyl)methylene]-*S*-[2-[(4-chlorophenyl)methylene]amino]ethyl]-L-cysteine, methyl ester.

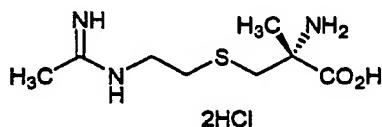
N-[4-Chlorophenyl)methylene]-*S*-[2-[(4-chlorophenyl)methylene]amino]ethyl]-2-methyl-D/L-cysteine, methyl ester.

The following examples are provided to illustrate the present invention and are not intended to limit the scope thereof. Those skilled in the art will readily understand that

5 known variations of the conditions and processes of the following preparative procedures can be used to prepare these compounds.

Example 1

10



S-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine, dihydrochloride

Example-1A) (2*R*,4*R*)-Methyl-2-tert-butyl-1,3-thiazoline-3-formyl-4-carboxylate

15 See Jeanguenat and Seebach, *J. Chem. Soc. Perkin Trans. 1*, 2291 (1991) and Pattenden *et al. Tetrahedron*, 49, 2131 (1993): (R)-cysteine methyl ester hydrochloride (8.58 g, 50 mmol), pivalaldehyde (8.61 g, 100 mmol), and triethylamine (5.57 g, 55 mmol) were refluxed in pentane (800 ml) with continuous removal of water using a Dean-Stark trap. The mixture was filtered and evaporated. The resultant thiazolidine (9.15 g, 45 mmol) and sodium

20 formate (3.37 g, 49.5 mmol) were stirred in formic acid (68 ml) and treated with acetic anhydride (13 mL, 138 mmol), dropwise over 1 hour at 0-5 °C. The solution was allowed to warm to RT and stir overnight. The solvents were evaporated and the residue was neutralized with aqueous 5% NaHCO₃, and extracted with ether (3X). The combined organic layers were dried (anhy. MgSO₄), filtered, and evaporated to give the title compound which

25 was crystallized from hexane-ether as white crystals (8.65 g) (80% overall, 8:1 mixture of conformers). ¹H NMR (CDCl₃) δ major conformer: 1.04 (s, 9H), 3.29 (d, 1H), 3.31 (d, 1H), 3.78 (s, 3H), 4.75 (s, 1H), 4.90 (t, 1H), 8.36 (s, 1H). MS m/z (electrospray) 232 (M+H)⁺ (100%), 204 (10) 164 (24).

Example-1B) (2R,4R)-Methyl-2-tert-butyl-1,3-thiazoline-3-formyl-4-methyl-4-carboxylate

To a solution of the product of **Example-1A**, (2R,4R)-Methyl-2-tert-butyl-1,3-thiazoline-3-formyl-4-carboxylate (8.65 g, 37.4 mmol), in anhydrous tetrahydrofuran (130 mL) under N₂ at -78 °C was added DMPU (25 mL) and the mixture stirred for 5 min. Lithium 5 bis(trimethylsilyl)amide, 1 M in tetrahydrofuran, (37.5 mL), was added, and the mixture stirred for 30 min. After methyl iodide (5.84 g, 41.1 mmol) was added, the mixture was held at -78 °C for 4 hr and then warmed to room temperature with continuous stirring. The solvents were evaporated *in vacuo* and brine and ethyl acetate was added. The aqueous phase was extracted 3x EtOAc, and the combined organic layers were washed with 10% 10 KHSO₄, water, and brine. They were then dried (anhy. MgSO₄), filtered, and stripped of all solvent under reduced pressure. Chromatography of the residual oil on silica with 1-10% EtOAc/hexane yielded the title compound (5.78 g, 63%, 2.4:1 mixture of conformers). ¹H NMR (CDCl₃) δ major conformer, 1.08 (s, 9H), 1.77 (s, 3H), 2.72 (d, 1H), 3.31 (d, 1H), 3.77 (s, 3H), 4.63 (s, 1H), 8.27 (s, 1H); minor conformer, 0.97 (s, 9H), 1.79 (s, 3H), 2.84 (d, 1H), 15 3.63 (d, 1H), 3.81 (s, 3H), 5.29 (s, 1H), 8.40 (s, 1H); MS m/z (electrospray) 246 (M+H)⁺ (100%), 188 (55) 160 (95). Retention time of 16.5 min on a Daicel Chemical Industries Chiracel OAS column, 10-40% IPA/hexane 0-25 min, >95% ee.

Example-1C) (2R) 2-Methyl-L-cysteine hydrochloride

20 The product of **Example-1B**, (2R,4R)-Methyl-2-tert-butyl-1,3-thiazoline-3-formyl-4-methyl-4-carboxylate, (5.7 g, 23.2 mmol) was stirred with 6N HCl (100mL) under N₂ and held at vigorous reflux for 2 days. The solution was cooled, washed with EtOAc and evaporated to yield the product (2R) 2-methyl-cysteine hydrochloride (3.79 g, 95%) as a light yellow powder. ¹H NMR (DMSO-d₆) δ 1.48 (s, 3H) 2.82 (t, 1H), 2.96 (bs, 2H), 8.48 (s, 25 3H). MS m/z (electrospray) 136 [M+H]⁺.

Example-1D) S-[2-[(1,1-dimethylethoxy)carbonyl]amino]ethyl]-2-methyl-L-cysteine trifluoroacetate

30 Sodium hydride (2.6 g, 60% in mineral oil, 65 mmol) was added to an oven-dried, vacuum-cooled RB flask, containing oxygen-free 1-methyl-2-pyrrolidinone (5 mL). The mixture was cooled to -10 °C and stirred under N₂. The product of **Example-1C**, 2-Methyl-

L-cysteine hydrochloride, (3.6 g, 21.0 mmol) dissolved in oxygen-free 1-methyl-2-pyrrolidinone (25 mL), was added in portions. After all H₂ evolution ceased, 2-[(1,1-dimethylethoxycarbonyl)-amino]ethyl bromide (4.94 g, 21 mmol) in oxygen-free 1-methyl-2-pyrrolidinone (15 mL) was added at -10 °C. The reaction was then stirred for 4 hr 5 allowing warming to room temperature. The solution was neutralized with 1 N HCl and the 1-methyl-2-pyrrolidinone was removed by evaporation *in vacuo*. Reverse-phase chromatography with 1-20% acetonitrile in 0.05% aqueous trifluoro acetic acid solution yielded the title compound (5.9 g), recovered by freeze-drying appropriate fractions. ¹H NMR (DMSO-d₆/D₂O) δ 1.31 (s, 9H), 1.39 (s, 3H), 2.55 (m, 2H), 2.78 (d, 1H), 3.04 (d, 1H), 10 3.06 (t, 2H). HRMS calc. for C₁₁H₂₂N₂O₄S: 279.1375 (M+H⁺), found 279.1379.

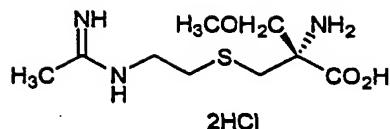
Example-1E) S-(2-aminoethyl)-2-methyl-L-cysteine hydrochloride

The product of **Example-1D**, S-[2-[(1,1-dimethylethoxy)carbonyl]amino]ethyl]-2-methyl-L-cysteine trifluoroacetate, (5.5 g, 14.0 mmol) was dissolved in 1 N HCl (100 mL) 15 and stirred at room temperature under nitrogen overnight. The solution was removed by freeze-drying to give the title S-(2-aminoethyl)-2-methyl-L-cysteine hydrochloride, ¹H NMR (DMSO-d₆/D₂O) δ 1.43 (s, 3H), 2.72 (m, 2H), 2.85 (d, 1H), 2.95 (t, 2H), 3.07 (d, 1H). m/z [M+H⁺] 179.

20 The product of **Example-1E**, was dissolved in H₂O, the pH adjusted to 10 with 1 N NaOH, and ethyl acetimidate hydrochloride (1.73 g, 14.0 mmol) was added. The reaction was stirred 15-30 min, the pH was raised to 10, and this process repeated 3 times. The pH was adjusted to 3 with HCl and the solution loaded onto a washed DOWEX 50WX4-200 column. The column was washed with H₂O and 0.25 M NH₄OH, followed by 0.5 M 25 NH₄OH. Fractions from the 0.5 M NH₄OH wash were immediately frozen, combined and freeze-dried to give an oil that was dissolved in 1N HCl and evaporated to give the title compound as a white solid (2.7 g). ¹H NMR (DMSO-d₆/D₂O) δ 1.17 (s, 3H), 2.08 (s, 3H), 2.52 (d, 1H), 2.68 (m, 2H), 2.94 (d, 1H), 3.23 (t, 2H). HRMS calc. for C₈H₁₈N₃O₂S: 220.1120 [M+H⁺], found 220.1133.

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Example 2



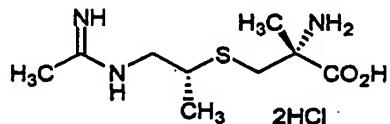
2-[[2-[(1-Iminoethyl)amino]ethyl]thio]methyl]-O-methyl-D-serine, dihydrochloride

5

The procedures and methods utilized in this example were identical to those of **Example 1** except that in step **Example-1B** methoxymethyl iodide was used instead of methyl iodide. These procedures yielded the title product as a white solid (2.7 g). ^1H NMR (D_2O) δ 2.06 (s, 3H), 2.70 (m, 3H), 3.05 (d, 1H), 3.23 (s, 3H), 3.32 (t, 2H), 3.46 (d, 1H), 3.62 (d, 1H).

10 HRMS calc. for $\text{C}_9\text{H}_{20}\text{N}_2\text{O}_3\text{S}$: 250.1225 $[\text{M}+\text{H}^+]$, found 250.1228.

Example 3



15

S-[(1R)-2-[(1-Iminoethyl)amino]-1-methylethyl]-2-methyl-L-cysteine, dihydrochloride

Example-3A) (S)-1-[(benzyloxycarbonyl)amino]-2-propanol

To a solution of *(S)*-1-amino-2-propanol (9.76 g, 130 mmol) in anhydrous benzene (60 mL) at 0 °C was added benzyl chloroformate (10.23 g, 60 mmol) in anhydrous benzene (120 mL) slowly, in portions, over a period of 20 min while vigorously stirring under an atmosphere of nitrogen. The mixture was stirred for 1 hour at 0 °C, then allowed to warm to room temperature and stirred for a further 2 hours. The mixture was washed with water (2X) and brine (2X) before the organic layer was dried over anhydrous MgSO_4 . Evaporation of all solvent gave the title product as an oil. ^1H NMR (CDCl_3) δ 1.22 (d, 3H), 2.40 (bs, 1H), 3.07 (m, 1H), 3.37 (m, 1H), 3.94 (m, 1H), 5.16 (s, 2H), 5.27 (m, 1H), 7.38 (m, 5H). MS m/z (electrospray) 232 $[\text{M}+23]^+$ (100%), 166 (96).

Example-3B) (S)-1-[(benzyloxycarbonyl)amino]-2-propanol tosylate

To a solution of the product of **Example-3A**, (S)-1-[(benzyloxycarbonyl)amino]-2-propanol, (9.74 g, 46.7 mmol) and triethylamine 7.27 g, 72 mmol) in methylene chloride (60 mL) at 0°C was added toluene sulfonyl chloride (9.15 g, 48 mmol) in methylene chloride (18 mL) slowly, in portions, over a period of 20 min while vigorously stirring under nitrogen. The mixture allowed to warm to room temperature and stirred for a further 36 hours under nitrogen. The organic layer was washed with 1N HCl, water, 5% NaHCO₃ solution, water and brine before it was dried over anhydrous MgSO₄. Evaporation of all solvent gave a white solid which was passed though a silica plug with ethyl acetate/hexane (1:4) to remove excess toluene sulfonyl chloride and then with ethyl acetate/hexane (1:3) to give the title product as white crystals. This material was recrystallized from ethyl acetate/hexane to give white needles (10.8 g). ¹H NMR (CDCl₃) δ 1.22 (d, 3H), 2.39 (s, 3H), 3.20 (m, 1H), 3.43 (dd, 1H), 4.66 (m, 1H), 5.02 (m, 1H), 5.04 (ABq, 2H), 7.34 (m, 7H), 7.77 (d, 2H). MS m/z (electrospray) 386 [M+23]⁺ (100%), 320 (66). The product was examined on a Regis Technologies Inc. Perkle Covalent (R,R) β-GEM1 HPLC column using mobile phase of isopropanol/hexane and a gradient of 10% isopropanol for 5 min, then 10 to 40% isopropanol over a period of 25 min, and using both UV and Laser Polarimetry detectors. Retention time major peak: 22.2 min, >98 % ee.

20

Example-3C) S-[(1R)-2-(Benzylloxycarbonylamino)-1-methylethyl]-2-methyl-L-cysteine trifluoroacetate

The product of **Example-1C**, 2-methyl-L-cysteine hydrochloride, (1 g, 6.5 mmol) was added to an oven dried, N₂ flushed RB flask, dissolved in oxygen-free 1-methyl-2-pyrrolidinone (5 mL), and the system was cooled to 0 °C. Sodium hydride (0.86 g, 60% in mineral oil, 20.1 mmol) was added and the mixture was stirred at 0 °C for 15 min. A solution of the product of **Example-3B**, (2S)-1-[(N-benzylloxycarbonyl)amino]-2-propanol tosylate (2.5 g, 7 mmol) dissolved in oxygen-free 1-methyl-2-pyrrolidinone (10 mL) was added over 10 min. After 15 min at 0 °C, the reaction mixture was stirred at room temperature for 4.5 hours. The solution was then acidified to pH 4 with 1N HCl and 1-methyl-2-pyrrolidinone was removed by evaporation *in vacuo*. Reverse phase

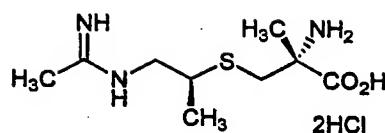
chromatography with 20-40 % acetonitrile in 0.05% aqueous trifluoro acetic acid solution yielded the title compound in (0.57g), recovered by freeze-drying. ^1H NMR (H_2O , 400 MHz) δ 1.0 (d, 3H), 1.4 (s, 3H), 2.6 (m, 2H), 2.8 (m, 1H), 3.1 (m, 2H), 3.6 (s, 1H), 5.0 (ABq, 2H), 7.3 (m, 5H). MS m/z (electrospray): 327 [M+H $^+$] (100%), 238 (20), 224 (10), and 100 (25).

Example-3D) S-[(1R)-2-Amino-1-methylethyl]-2-methyl-L-cysteine hydrochloride

The product of **Example-3C**, *S*-[(1R)-2-(Benzylloxycarbonylamino)-1-methylethyl]-2-methyl-L-cysteine trifluoroacetate, (0.5 g, 1.14 mmol) was dissolved in 6N HCl and refluxed for 1.5 hour. The mixture was then cooled to room temperature and extracted with EtOAc. The aqueous layer was concentrated *in vacuo* to give the title product, (2R, 5R)-*S*- (1-amino-2-propyl)-2-methyl-cysteine hydrochloride (0.29 g), which was used without further purification. ^1H NMR (H_2O , 400 MHz) δ 1.2 (m, 3H), 1.4 (m, 3H), 2.7 (m, 1H), 2.8-3.2 (m, 2H), 3.4 (m, 1H). (some doubling of peaks due to rotameric forms). MS m/z (electrospray): 193 [M+H $^+$] (61%), 176 (53), 142 (34), 134 (100), and 102 (10).

The product of **Example-3D**, *S*-[(1R)-2-Amino-1-methylethyl]-2-methyl-L-cysteine hydrochloride, (0.2 g, 0.76 mmol) was dissolved in 2 mL of H_2O , the pH was adjusted to 10.0 with 1N NaOH, and ethyl acetimidate hydrochloride (0.38 g, 3 mmol) was added in four portions over 10 minutes, adjusting the pH to 10.0 with 1N NaOH as necessary. After 1h, the pH was adjusted to 3 with 1N HCl. The solution was loaded onto a water-washed DOWEX 50WX4-200 column. The column was washed with H_2O and 0.5N NH₄OH. The basic fractions were pooled and concentrated to dryness *in vacuo*. The residue was acidified with 1N HCl and concentrated to the **Example 3** title product, (49 mg). ^1H NMR (H_2O , 400 MHz) δ 1.3-1.0 (m, 3H), 1.5 (m, 3H), 2.1-1.8 (m, 3H), 3.4-2.6 (m, 5H), 3.6 (m, 1H) (rotamers observed). MS m/z (electrospray): 234 [M+H $^+$] (100%), 176 (10), and 134 (10).

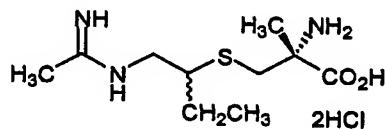
Example 4



S-[(1S)-2-[(1-Iminoethyl)amino]-1-methylethyl]-2-methyl-L-cysteine, dihydrochloride

5 The procedures and methods employed here were identical to those of **Example 3**, except that in step **Example-3A** (*R*)-1-amino-2-propanol was used instead of (*S*)-1-amino-2-propanol to give the title material, *S*-[(1*S*)-2-[(1-Iminoethyl)amino]-1-methylethyl]-2-methyl-L-cysteine hydrochloride. ^1H NMR (H_2O , 400 MHz) δ 3.6 (m, 1H), 3.4-2.6 (m, 5H), 2.1-1.8 (m, 3H), 1.5 (m, 3H), and 1.3-1.0 (m, 3H). HRMS calc for $\text{C}_9\text{H}_{19}\text{N}_3\text{O}_2\text{S}$ [M+H $^+$]: 10 234.1276. Found: 234.1286.

Example 5

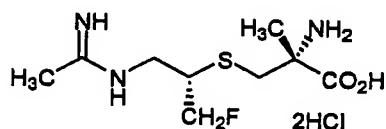


15

S-[(1*R/S*)-2-[(1-Iminoethyl)amino]-1-ethylethyl]-2-methyl-L-cysteine, dihydrochloride

The procedures and methods utilized in this synthesis are identical to those of **Example 3**, except that in step **Example-3A** (*R/S*)-1-amino-2-butanol is used instead of (*S*)-1-amino-2-propanol.

Example 6

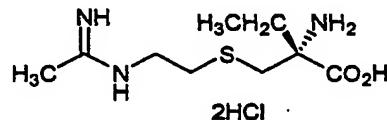


25

***S*-[*(1S*)-2-[(1-iminoethyl)amino]-1-(fluoromethyl)ethyl]-2-methyl-L-cysteine,
dihydrochloride**

A sample of 2-[(2*R*)-Oxiranylmethyl]-1*H*-isoindole-1,3-dione (G. Alexander et al. Tetrahedron Asymmetry, 7, 1641-8, 1996) is treated with potassium hydrogen difluoride to give 2-[(2*R*)-3-fluoro-2-hydroxypropyl]-1*H*-isoindole-1,3-dione in the presence of catalyst n*Bu*₄NH₂F₃. The procedures and methods used in this synthesis are identical to those of Example 3, except that in step Example-3B 2-[(2*R*)-3-fluoro-2-hydroxypropyl]-1*H*-isoindole-1,3-dione is used instead of (*S*)-1-[(benzyloxycarbonyl)amino]-2-propanol to produce the title product.

Example 7



15

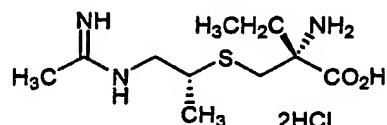
***S*-[2-[(1-Iminoethyl)amino]ethyl]-2-ethyl-L-cysteine, dihydrochloride**

The procedures and methods used in this synthesis were the same as those used in Example 1 except that ethyl triflate was used in Example-1B instead of methyl iodide.

20 Reverse phase chromatography, using a gradient of 10-40% acetonitrile in water, was used to purify the title product (20% yield). ¹H NMR (D₂O) δ 0.83 (t, 3H), 1.80 (m, 2H), 2.08 (s, 3H), 2.68 (m, 1H), 2.78 (m, 1H), 2.83 (m, 1H), 3.11 (m, 1H), 3.36 (t, 2H). HRMS calc. for C₉H₂₀N₃O₂S: 234.1276 [M+H⁺], found 234.1284.

25

Example 8

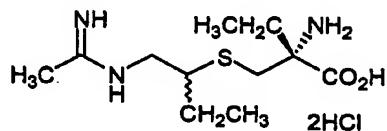


S-[(1*R*)-2-[(1-Iminoethyl)amino]-1-methylethyl]-2-ethyl-L-cysteine, dihydrochloride

The procedures and methods employed in this synthesis are the same as those used in
 5 **Example 1** except that ethyl triflate is used in **Example 1B** instead of methyl iodide. The 2-ethyl-L-cysteine hydrochloride thus prepared is treated as described in the procedures and methods of **Examples 3C - 3E** to give the title compound.

Example 9

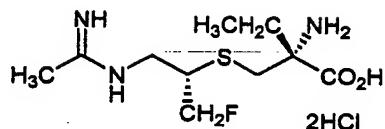
10

***S-[(1*R/S*)-2-[(1-Iminoethyl)amino]-1-ethylethyl]-2-ethyl-L-cysteine, dihydrochloride***

15 The procedures and methods utilized in this synthesis are the same as those used in
Example 5 except that 2-ethyl-L-cysteine hydrochloride (prepared in **Example 7**) is used instead of 2-methyl-L-cysteine hydrochloride to give the title compound.

Example 10

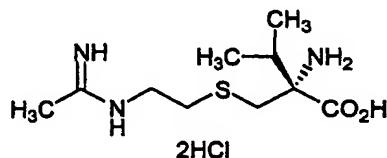
20

***S-[(1*S*)-2-[(1-iminoethyl)amino]-1-fluoromethylethyl]-2-ethyl-L-cysteine, dihydrochloride***

25

The procedures and methods utilized in this synthesis are the same as those used in **Example 6**, except that (2*R*)-2-ethyl cysteine hydrochloride (prepared in **Example 7**) is used instead of (2*R*)-2-methyl cysteine hydrochloride, to give the title compound.

5

Example 11**2-[[[2-(1-Iminoethyl)amino]ethyl]thio]methyl-D-valine, dihydrochloride**

10

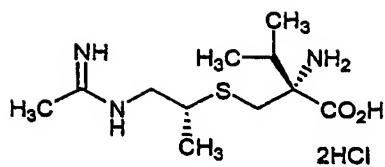
Example-11a) Isopropyl triflate

Silver triflate (25.25 g, 98.3 mmol) stirred in diethyl ether (300 mL) under nitrogen was treated with isopropyl iodide (16.54 g, 98.5 mmol) in ether (200 mL) over 15 minutes. The mixture was stirred for 10 minutes and then filtered. The filtrate was distilled at reduced pressure. The distillate was redistilled at atmospheric pressure to remove the majority of the diethyl ether, leaving a mixture of the title isopropyl triflate-diethyl ether (84:16 by weight) (15.64 g, 70% corrected) as a colorless liquid. ^1H NMR (CDCl_3 , 400 MHz) δ 1.52 (d, 6H), 5.21 (septet, 1H).

20

The procedures and methods utilized here were the same as those used in **Example 1** except that isopropyl triflate replaced methyl iodide in **Example-1B**. The crude title product was purified by reversed phase chromatography using a gradient elution of 10-40% acetonitrile in water. ^1H NMR (H_2O , 400 MHz) δ 0.94 (dd, 6H), 2.04 (septet, 1H), 2.10 (s, 3H), 2.65, 2.80 (d m, 2H), 2.85, 3.10 (dd, 2H), 3.37 (t, 2H). HRMS calc. for $\text{C}_{10}\text{H}_{22}\text{N}_3\text{O}_2\text{S}$:

25 248.1433 [M+H $^+$], found 248.1450.**Example 12**



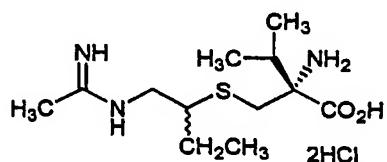
**2-[[[(1*R*)-2-[(1-Iminoethyl)amino]-1-methylethyl]thio]methyl]-D-valine,
dihydrochloride**

5

The procedures and methods used in this synthesis are the same as those used in **Example 3** except that isopropyl triflate (prepared in **Example-11A**) is used instead of methyl iodide to give the title compound.

10

Example 13



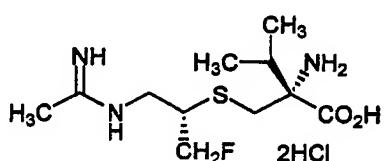
15

**2-[[[(1*R*/*S*)-2-[(1-Iminoethyl)amino]-1-ethylethyl]thio]methyl]-D-valine,
dihydrochloride**

The procedures and methods used in this synthesis are the same as those used in **Example 5**, except that (2*R*)-2-isopropyl triflate (prepared in **Example-11A**) is used instead of methyl iodide, to give the title compound.

20

Example 14

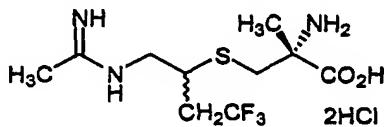


**2-[[[(1*S*)-2-[(1-Iminoethyl)amino]-1-fluoromethylethyl]thio]methyl]-D-valine,
dihydrochloride**

The procedures and methods used in this synthesis are the same as those used in **Example 5** 6, except that (2*R*)-2-isopropyl triflate (prepared in **Example-11A**) is used instead of methyl iodide, to give the title compound.

Example 15

10

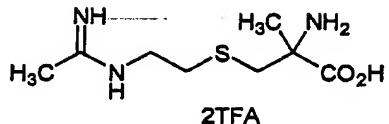


**S-[(*R/S*)-2-[(1-iminoethyl)amino]-1-(trifluoromethyl)ethyl]-2-methyl-L-cysteine,
dihydrochloride**

15 *t*-Butyl-N-(2-oxoethyl)carbamate is treated with 1,1,1-trifluoroethyl magnesium bromide to give (*R/S*)-1-[(1,1-dimethylethoxycarbonyl)]amino-4,4,4-trifluoro-2-butanol. The procedures and methods used in this synthesis are the same as those listed in **Example 3**, except that (*R/S*)-1-[(1,1-dimethylethoxy)carbonyl]amino-4,4,4-trifluoro-2-butanol is used instead of (*S*)-1-[(benzyloxycarbonyl)amino]-2-propanol, to give the title compound.

20

Example 16



25 **S-[2-(1-Iminoethylamino)ethyl]-2-methyl-(D/L)-cysteine, bistrifluoroacetate**

Example-16A) S-(2-aminoethyl)-L-cysteine, methyl ester

A 10 g (50 mmol) sample of *S*-(2-aminoethyl)-L-cysteine was dissolved in 400 mL of 30 methanol. Into this cooled solution was bubbled in anhydrous HCl for 30 minutes. After

stirring at room temperature overnight, the solution was concentrated to afford 12.7 g of the title compound.

Example-16B) *N*-(4-chlorophenyl)methylene]-S-[2-[(4-

5 **chlorophenyl)methylene]amino]ethyl]-L-cysteine, methyl ester**

A 12.7 g (50 mmol) sample of the product of **Example-16A, S-(2-aminoethyl)-L-cysteine methyl ester**, was dissolved in acetonitrile. To this solution was added 12.2 g (100 mmol) of anhydrous $MgSO_4$, 14g (100 mmol) of 4-chlorobenzaldehyde and 100 mmol of triethylamine. This mixture was stirred for 12 hours, concentrated to a small volume and 10 diluted with 500 mL of ethyl acetate. The organic solution was washed successively with (0.1%) $NaHCO_3$, (2N) $NaOH$, and brine solution. The organic was dried (anhy. $MgSO_4$), filtered and concentrated to afford 7.5g of the title compound. $[M + H^+] = 179$.

Example-16C) *N*-(4-chlorophenyl)methylene]-S-[2-[(4-

15 **chlorophenyl)methylene]amino]ethyl]-2-methyl-D/L-cysteine methyl ester**

A sample of the product of **Example-16B, N-(4-chlorophenyl)methylene]-S-[2-[(4-chlorophenyl)methylene]amino]ethyl]-L-cysteine methyl ester** (7.5 g, 17 mmol), in anhydrous THF was treated with 17 mmol of sodium bis(trimethylsilyl)amide at -78 °C under nitrogen, followed by 2.4g (17mmol) of methyl iodide. The solution was held at -78 20 °C for 4 hr and then warmed to room temperature with continuous stirring. The solvents were evaporated *in vacuo* and brine and ethyl acetate was added. The aqueous phase was extracted 3x EtOAc, and the combined organic layers were washed with 10% $KHSO_4$, water, and brine before it was dried (anhy. $MgSO_4$), filtered, and evaporated to afford the title compound.

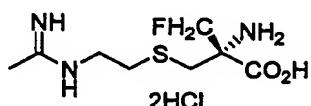
25

Example-16D) *S*-(2-aminoethyl)-2-methyl-D/L-cysteine, hydrochloride

A sample of the product of **Example-16C, N-(4-chlorophenyl)methylene]-S-[2-[(4-chlorophenyl)methylene]amino]ethyl]-2-methyl-D/L-cysteine methyl ester** (4.37 g, 10 mmol), was stirred and heated (60 °C) with 2N HCl overnight and the solution washed (3X) 30 with ethyl acetate. The aqueous solution was freeze-dried to give the title compound.

A sample of the product of **Example 16D**, *S*-(2-aminoethyl)-2-methyl-D/L-cysteine dihydrochloride (2.5 g (10 mmol), was dissolved in H₂O and the pH was adjusted to 10 with 1 N NaOH. Ethyl acetimidate hydrochloride (1.24 g, 10.0 mmol) was then added to the reaction mixture. The reaction was stirred 15-30 min, the pH was raised to 10, and this process repeated 3 times. The pH was reduced to 4 with HCl solution and the solution evaporated. The residue was purified on reverse phase HPLC with H₂O containing 0.05% trifluoroacetic acid as the mobile phase to afford the **Example 16** title product. M + H = 220.

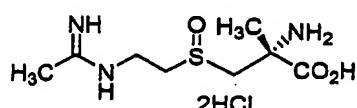
10

Example 17**S-[2-[(1-Iminoethyl)amino]ethyl]-2-fluoromethyl-L-cysteine, dihydrochloride**

15

Epibromohydrin is treated with HF-pyridine to give the dihalogenated alcohol, which is oxidized with K₂Cr₂O₇ to give the 1-bromo-3-fluoroacetone. This product is treated with (1,1-dimethylethoxy)-N-(2-sulfanyethyl)carboxamide in the presence of NaOH to give (1,1-dimethylethoxy)-N-[2-(3-fluoro-2-oxopropylthio)ethyl]carboxamide. This is cyclized to the racemic hydantoin by NaCN and (NH₄)₂CO₃ in refluxing ethanol and the enantiomers separated by chiral chromatography. The S-enantiomer is treated with hot 48% HBr solution to afford *S*-(2-aminoethyl)-2-fluoromethyl-L-cysteine dihydrochloride, which is converted to the title compound by treatment with ethyl acetimidate in the presence of base.

25

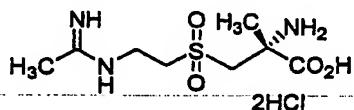
Example 18

**(2*R*)-2-Amino-3[[2-[(1-iminoethyl)amino]ethyl]sulfinyl]-2-methylpropanoic acid,
dihydrochloride**

A solution of *S*-[2-[(1-iminoethyl)amino]ethyl]-2-methyl-L-cysteine, dihydrochloride (Example 1, 0.2g, 0.73 mmol) in 3 mL of water was stirred and cooled to 0 °C and a solution of 3% H₂O₂ (0.8 mL, 0.73 mmol) in formic acid (0.4 mL, 0.73 mmol) was added in 0.3 mL portions. The cold bath was removed and the reaction mixture was stirred at room temperature for 48 hours. The solution was concentrated *in vacuo*, diluted with of water (10 mL) and concentrated again to give the crude sulfone. This residue was chromatographed (C-18 reverse phase, with mobile phase H₂O containing 0.05% trifluoroacetic acid) to give the pure sulfone. The sulfone was treated with 1M HCl (10 mL) and concentrated *in vacuo* to give 140 mg of a mixture of 2 diastereomers of the title compound as a colorless oil of the HCl salts. ¹H NMR (300 MHz, D₂O) δ 1.5 (s, 2H), 1.6 (s, 1H), 2.0 (s, 3H), 3.1 (m, 2H), 3.3 (m, 2H) 3.6 (m, 2H). HRMS calc. for C₈H₁₈N₃O₃S: 236.1069 [M+H⁺], found: 236.1024.

15

Example 19

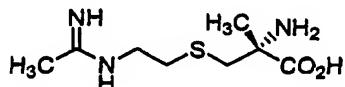


20 **(2*R*)-2-Amino-3[[2-[(1-iminoethyl)amino]ethyl]sulfonyl]-2-methylpropanoic acid
dihydrochloride**

A solution of *S*-[2-[(1-iminoethyl)amino]ethyl]-2-methyl-L-cysteine dihydrochloride, the product of Example 1, (0.15 g, 0.54 mmol) in 2 mL of water was cooled to 0 °C and a solution of 3% H₂O₂ (1.6 mL, 1.46 mmol) in formic acid (0.8mL, 14.6 mmol) was added. The cold bath was removed and the reaction mixture was stirred at room temperature for 18 hours. The solution was concentrated *in vacuo*, diluted with 10 mL of water and concentrated again to give the crude sulfoxide. The residue was diluted with 4 mL of water and was adjusted to pH 9 with 2.5 N NaOH. Acetone (5 mL) was added, followed by Boc₂O (0.2 g), and the reaction was stirred for 48 h at room temperature. The reaction mixture was

adjusted to pH 6 with 1M HCl and was concentrated *in vacuo*. This residue was chromatographed (C-18 reverse phase; 40 to 50% ACN: H₂O, 0.05% TFA) to give the pure Boc protected material. The fractions were concentrated *in vacuo* and the residue was treated with 1N HCl (3 mL) for 1h. The solution was concentrated to give 30 mg of the title 5 compound as colorless oil. ¹H NMR (400 MHz, D₂O) δ 4.0 (d, 1H), 3.7 (d, 1H), 3.6 (t, 2H), 3.5 (t, 2H), 2.1 (s, 3H), and 1.5 (s, 3H) ppm. HRMS calc. for C₈H₁₈N₃O₄S: 252.1018 [M + H⁺], found: 252.0992.

10 **Example 20**

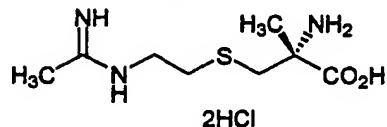


15 **S-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine**

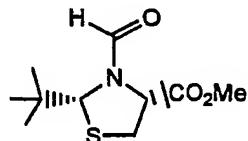
15 DOWEX 50WX4-200 (250 g) in a glass chromatography column (38 X 560 mm) was washed with water until the eluent was at pH 6. An solution of the product of Example 1, S-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine dihydrochloride (6 g) dissolved in water was placed on the column, which was washed with water until the pH returned to 6. The column was then washed with 0.07 M NH₄OH (flow rate ~15mL/min) and the basic fractions 20 were immediately placed in a dry ice / acetone bath. The fractions were pooled and concentrated to dryness by lyophilization to give the title compound. ¹H NMR (400MHz, DMSO-d₆) δ 3.4 (m, 1H), 3.3 (m, 1H), 3.0 (d, 1H), 2.7 (m, 1H), 2.4 (m, 1H), 2.3 (d, 1H), 2.1 (s, 3H), and 1.1 (s, 3H).

25 Analysis calculated for C₈H₁₇N₃O₂S + 0.6 H₂O: C 41.76, H 7.97, N 18.26; found C 41.43, H 7.47, N 17.96, Cl trace.

30 **Example 21**

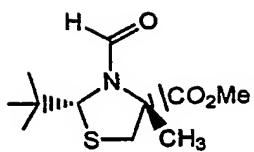


S-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine, dihydrochloride

Example-21A) (2*R*,4*R*)-Methyl-2-tert-butyl-1,3-thiazoline-3-formyl-4-carboxylate

5 The title material was prepared according to J. Chem. Soc. Perkin Trans. 1991 p 2291 and Tetrahedron 1993 p 2131. To a 2 L RB flask fitted with a reflux condenser, a Dean-Stark trap, an overhead stirrer and a thermocouple was added pivalaldehyde (23.7 g, 0.275 mole) dissolved in 700 mL of toluene. Agitation was started and L-cysteine hydrochloride methyl ester (45 g, 0.262 mole) was added to the stirring solution. Triethylamine (29.2 g, 0.288 moles) was then added to the batch in a stream over a few minutes. The reaction mixture was heated to reflux and water was removed. The batch was heated for a total of 3h, cooled and filtered. The salt cake was washed with 250mL of fresh toluene and the wash was combined. Formic acid (24.1 g, 0.524 moles) and solid sodium formate (19.6 g, 0.288 moles) were then added and the resulting suspension was cooled to -5 °C. Acetic anhydride (53.5 g, 0.524 moles) was carefully added to the mixture keeping the batch temperature below 5 °C. After the addition, the reaction was allowed to warm to room temperature and stirring continued for 18h during which time the product precipitated. The crude product was filtered and redissolved in 400 mL of EtOAc and filtered to remove insoluble sodium salts. The organic solution was then neutralized with 200 mL of saturated sodium bicarbonate solution and the final aqueous layer was near pH 7. The organic layer was separated and the aqueous layer was extracted with ethyl acetate. The organic layers were combined and concentrated to give the crude product (60.2 g) as a viscous oil which slowly crystallized to a white solid. The solid was washed with cyclohexane containing 4% of 2-propanol to give 41.01 g of the title product in >99.5% purity and 67.8% yield as determined by GC. The desired title product cis isomer was present in over 98%.

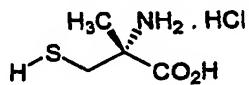
Example-21B) (2*R*,4*R*)-Methyl-2-tert-butyl-1,3-thiazoline-3-formyl-4-methyl-4-carboxylate



Anhydrous lithium chloride (43.0 g, 0.102 moles) was mixed with 300 mL of dimethoxyethane and 500 mL of THF until a clear solution was obtained. A THF solution of the product of **Example-21A** (50.0 g, 0.216 moles) was added and cooled to -65 °C under a nitrogen atmosphere. Iodomethane (45.0 g, 0.316 moles) diluted with 45 mL of THF was added followed by 230 mL of 1.0M THF solution of lithium bis-trimethylsilylamide. The reaction was stirred at -65 °C for 10 h. The batch was quenched with 30 g of acetic acid in 600 mL water and extracted with 500 mL of ethyl acetate. The organic layer was washed with saturated sodium bicarbonate and concentrated to give 51.22 g (96%) of the title product as a light brown solid.

Example-21C) (2R) 2-Methyl-L-cysteine hydrochloride

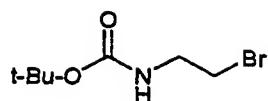
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A sample of the product of **Example-21B** (20 g, 83 mmoles) was placed in a round bottom flask equipped with an overhead stirrer and reflux condenser. To this solid was added 100 mL of concentrated hydrochloric acid. The reaction was slowly heated to 95 °C for 7 days. The reaction was then treated with 250 mL of toluene to remove non-polar organic impurities. The aqueous solution was then concentrated. The crude title product was obtained as an orange resin weighing 14 g. The resin was powdered up under ethylether / methylene chloride and filtered to give 13 g of the title material as a pale brown hygroscopic powder. ¹H NMR (D₂O) δ 4.70 (s, HDO exchange), 3.08 (d, 1H), 2.80 (d, 1H), 1.48 (s, 3H).

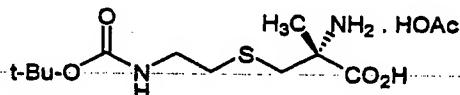
25

Example-21D) 2-[(1,1-dimethylethoxycarbonyl)-amino]ethyl bromide



A 5L RB flask equipped with overhead stirrer, thermocouple and nitrogen inlet was charged with 3 liters of ethyl acetate and agitation started. To this was added di-*tert*-butyldicarbonate (545 g, 2.50 moles) and 2-bromoethylamine hydrobromide (553.7 g, 2.70 moles) under a nitrogen blanket. The batch was cooled to 5 °C in an ice bath and N-methylmorpholine, (273 g, 2.70 moles) was added dropwise over ca. 0.5 h. After the addition was complete, the batch was allowed to stir overnight and warm to ambient temperature. After 16h, the batch was quenched by adding 1.5L of DI water. The organic layer was washed with dilute HCl, sodium bicarbonate solution followed by brine. The dried organic solution had solvents removed to give an oil that froze to a light yellow solid. A total of 496 g (88% yield) of the title product was obtained in approx. 96 % purity.

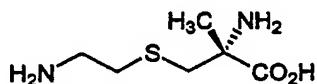
Example-21E) *S*-[2-[(1,1-dimethylethoxy)carbonyl]amino]ethyl]-2-methyl-L-cysteine acetate



A 3 L flask, charged with 435 g of methanol, was equipped with an overhead stirrer, a thermocouple, and maintained under a N₂ purge. To the reaction flask was added 150.0 g (0.804 moles) of the product of **Example-21C** carefully with stirring to dissolve. A solution of KOH prepared by dissolving 154.7 g of solid KOH in 840 mL of degassed methanol was added to the reaction solution dropwise keeping the temperature between 20-30 °C. The product of **Example-21C** (180.2 g, 0.804 moles) was dissolved in 375 mL of methanol and this solution was added dropwise to the cold reaction mixture over 1 hour at 10-12 °C. When the reaction was complete, the batch had its pH adjusted to pH 5. The reaction mixture was then filtered through a pad of Celite and the filtrate was concentrated to yield 454 g of a tan solid product. This title product was slurried with ethyl acetate to give an off-white solid

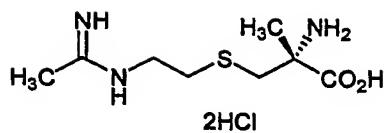
weighing 299 g. The crude title product solid was carried on to the next step without further purification. ^1H NMR as acetate (D_2O) δ 4.68 (s, D_2O exchange), 3.12 (m, 3H), 2.68 (m, 3H), 1.83 (s, 3H), 1.42 (s, 3H), 1.32 (s, 9H).

5 **Example-21F) S-(2-aminoethyl)-2-methyl-L-cysteine**



To a round bottom flask equipped with an overhead stirrer and nitrogen purge was added 10 150 mL of 37% hydrochloric acid. The agitator was started and 150 mL of water was added to the vessel followed by 173 g of crude product of Example-21E. The reaction was stirred for two hours and the clear brown solution batch was concentrated to give the crude di-HCl salt of the title product as a brown syrup (ca.157 g). This was redissolved in 200 mL of water and decolorized with charcoal. The solution was passed through a Dowex resin column 15 and the neutral title product was eluted with aqueous ammonium hydroxide to yield 64% of this material with approx. 94% purity. ^1H NMR (D_2O , 300MHz) δ 4.68(s, D_2O exchange), 2.9(m, 3H), 2.6(m, 3H), 1.20(s, 3H).

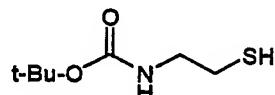
20 Polymer bound triazabicyclo[4.4.0] dec-5-ene resin (Fluka), 38 g, was suspended in 160 mL of ethanol in a round bottom flask. The amino acid product of Example-21F (7.5 g) in 40 mL of ethanol was added to the stirring resin slurry. Ethyl acetimidate, 6.5 g (53 mmoles) was added portion-wise to the reaction. The reaction was stirred under nitrogen for 16h. The resin was filtered and washed on the filter with 100 mL of ethanol containing 10mL of conc. HCl. The combined filtrate was concentrated to give 12 g of crude title 25 product as a pale brown viscous semisolid. The yield was approximately 60-70% and the title product showed 90% purity. The title product was further purified by reverse phase chromatography. ^1H NMR (D_2O) δ 4.74 (s, D_2O exchange), 3.37 (t, 2H), 3.08 (d, 1H), 2.93 (d, 1H), 2.74 (m, 2H), 2.06 (s, 3H), 1.48 (s, 3H).



S-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine, dihydrochloride

Example-22A) N-Boc-cysteamine

5

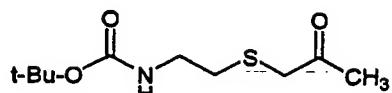


A 3L 4-neck RB flask was purged with nitrogen for 20 min and then charged sequentially with 2-aminoethanethiol hydrochloride (113.6 g, 1 mol), di-tert-butyl-dicarbonate (218.3 g, 1 mol) and 500 mL of toluene. The mixture was cooled with an ice-water bath and purged with nitrogen for 10 min. Sodium hydroxide (2.5N, 880 mL, 2.2 mol) was added to the stirring mixture in about 1.5 h at between 0 and 11 °C. After the addition of sodium hydroxide was complete, the cooling bath was removed and the resulting reaction mixture was allowed to warm up to room temperature and stirred at ambient temperature overnight.

10 This provided a solution of the title product.

15

Example-22B)



20

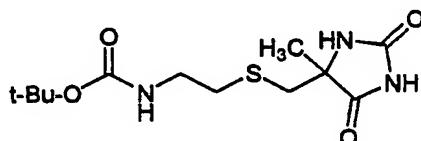
The product solution of Example-22A was cooled with an ice-water bath. A sample of chloroacetone (101.8 g, 1.1 mol) was added to the vigorously stirred reaction mixture over about 50 min at between 8 and 11 °C. After the addition of chloroacetone was completed, the cooling bath was removed and the resulting reaction mixture was allowed to stir at room temperature overnight. The toluene layer was separated, washed with water (250 mL) and concentrated on a rotary evaporator at 85 °C under house vacuum followed by high vacuum

25

to give the crude titled compound (225.7 g, 96.7%). ^1H NMR (CDCl_3 , 400 MHz) δ 4.95 (bs, 1H), 3.20 (m, 4H), 2.54 (t, 2H), 2.20 (s, 3H), 1.35 (s, 9H).

Example-22C) [2-[[[4-Methyl-2,5-dioxo-4-imidazolidinyl)methyl]thio]ethyl]carbamic acid,

5 1,1-dimethylethyl ester



To a 3L 4-neck RB flask equipped with an overhead stirrer, a thermocouple and a condenser connected to an empty flask and a caustic trap, was added the product of Example-22B (70 g, 0.3 mol), absolute ethanol (80 mL), sodium cyanide (19.1 g, 0.39 mol), ammonium carbonate (43.3 g, 0.45 mol) and water (720 mL) in this order. The 4th neck was closed with a stopper. The resulting reaction mixture was heated at between 67 and 68 °C for 6 h. Subsequently, the almost clear brown solution was cooled to room temperature.

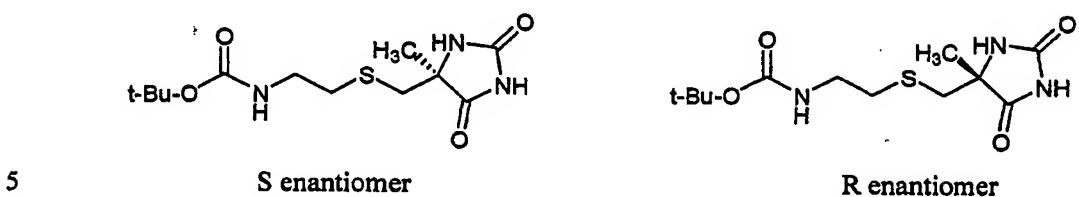
10 Upon cooling, solid began to form and the heterogeneous mixture was stirred at room temperature overnight. The reaction mixture was then acidified with 12% hydrochloric acid to pH 2 in about 1 h at between -2 and 2 °C. The cold reaction mixture was stirred at pH 2 for additional 30 min and then filtered. The flask was rinsed with distilled water (2 x 250 mL) and each rinse was used to wash the solid cake. The solid was again washed with

15 distilled water (2 x 250 mL) and then air-dried for 4 days. The dry solid was triturated with 200 mL of toluene for 0.5 h. The slurry was filtered. The solid was rinsed sequentially with toluene (50 mL) and 1:4 ratio of toluene/hexane (100 mL) and then air-dried at room temperature overnight to give 83.1% yield of the titled compound, m.p. 134-136 °C. ^1H NMR ($\text{DMSO}_{\text{d}6}$, 400 MHz) δ 10.62 (s, 1H), 7.85 (s, 1H), 6.83 (m, 0.9H), 6.48 (bs, 0.1H), 3.29 (s, 2H), 2.99 (m, 2H), 2.71 (s, 2H), 2.95 (m, 2H), 1.32 (s, 9H), 1.24 (s, 3H); ^{13}C NMR ($\text{DMSO}_{\text{d}6}$, 400 MHz), δ 178.1, 157.1, 156.1, 78.4, 63.7, 40.7, 39.4, 33.2, 28.9, 23.8.

20 Analysis Calcd for $\text{C}_{12}\text{H}_{21}\text{N}_3\text{O}_4\text{S}$: C, 47.51; H, 6.98; N, 13.85; S, 10.57. Found: C, 47.76; H, 6.88; N, 13.77; S, 10.75.

25

Example-22D) R and S-[2-[(4-Methyl-2,5-dioxo-4-imidazolidinyl)methyl]thio]ethyl]carbamic acid, 1,1-dimethylethyl ester



The reaction product of **Example-22C** was separated into its R and S enantiomers on a Chiralpak® AD column eluting with methanol. The S isomer was the first eluting isomer followed by its R enantiomer. Both isomers were used in subsequent transformations.

10 S enantiomer:

$[\alpha]$ in MeOH at 25 °C = +43.0 (365 nm). ^1H NMR: (400mHz, CD₃OD) δ 1.49 (s, 9H), 2.05 (s, 3H), 2.65 (t, 2H), 2.9 (q, 2H, d), 3.20 (m, 2H). IR: λcm^{-1} = 1772, 1709.

Analysis calculated for $C_{12}H_{21}N_3O_4S$ (formula weight = 303.38): C 47.51, H 6.98, N 13.85.
 Found: C 47.39, H 6.62, N 13.83. M+H = 304.

15

Renantiomer:

$[\alpha]$ in MeOH at 25 °C = -46.3 (365 nm). ^1H NMR: (400MHz, CD_3OD) δ 1.48 (s, 9H), 2.05 (s, 3H), 2.65 (t, 2H), 2.85 (q, 2H, d), 3.18 (m, 2H). IR: λcm^{-1} = 1770, 1711.

Analysis calculated for C₁₂H₁₀N₂O₂S (formula weight = 303.38): C 47.51, H 6.98, N 13.85.

20 Found: C 48.15, H 7.04, N 14.37. M+H = 304.

Example-22E) S-(2-aminoethyl)-2-methyl-L-cysteine



Acid hydrolysis method:

A 500 mL three-necked round bottom flask equipped with a distillation condenser was charged with the R-isomer product of **Example-22D** (45.8 g, 150.9 mmol) and treated portion wise with 48% aq. HBr (160 mL) at room temperature with stirring. After the gas evolution ceased, the mixture was heated with a heating mantle until the pot temperature 5 reached to 126 °C while the volatile t-butyl bromide (bp 72-74 °C) followed by a small amount of aq. HBr (approx. 15 mL) were distilled off. The distillation condenser was replaced with a reflux condenser and the mixture was heated at reflux for 30 hours. The solution was concentrated and the residue was dissolved in water (250 mL) and loaded on to a Dowex® 50WX4-200 ion-exchange resin (8.5 x 11 cm) and eluted with water (2L) 10 followed by dilute aqueous ammonium hydroxide (30 mL of 28-30% ammonium hydroxide diluted to 1000 mL with water, 3L). Fractions containing the desired product were combined, concentrated, and dried under vacuum at 75-80 °C for two hours to give 22.1 g (82%) of the title product, S-(2-aminoethyl)-2-methyl-L-cysteine, as a white solid. Proton and C-13 NMR spectra are consistent with the title product. Mp 157 °C. ¹H NMR (400 15 MHz, D₂O) δ 1.19 (3H, s), 2.53 (1H, d, *J* = 13.6 Hz), 2.57 – 2.72 (2H, m), 2.92 (1H, d, *J* = 13.6 Hz), 2.92 (2H, t, *J* = 6.8 Hz); ¹³C NMR (100 MHz, D₂O) δ 24.7, 31.3, 38.9, 40.9, 59.6, 180.7.

Analysis Calcd for C₆H₁₄N₂O₂S + 0.1 H₂O: C, 40.02; H, 7.95; N, 15.56; S, 17.81. Found: C, 39.93; H, 7.98; N, 15.38; S, 17.70.

20

Base hydrolysis method:

To a stainless steel autoclave equipped with agitation was added 24.2 g (0.08 moles) of the R-isomer product of **Example-22D**. After purging the apparatus with nitrogen, 128 g (0.32 moles) of 10% caustic was added generating a solution. The autoclave was sealed and 25 heated to 120 °C for 30 hours. After cooling to room temperature, the autoclave was vented to give 142 ml (151 g) of an aqueous solution of the sodium salt of the title product. ¹H NMR (sample acidified with HCl and diluted with D₂O, 400 MHz): δ 1.47 (s, 3H), 2.75 (m, 2 H), 2.90 (d, 1H, *J* = 14.8 Hz), 3.06 (t, 2H, *J* = 6.4 Hz), 3.14 (d, 1H, *J* = 14.8 Hz). ¹³C NMR (sample acidified with HCl and diluted with D₂O, 100 MHz): δ 172.9, 60.8, 39.1, 39.0, 30.4, 30 22.2. MS (MS/CI-LC) M+1 179.

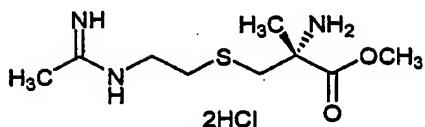
DBU (218 L; 1.46 mmol) and ethyl acetimidate hydrochloride (171 mg; 1.34 mmol) were dissolved in ethanol (6 mL) in a 25 mL, one-necked, round-bottomed flask at room temperature (~20°C). The title product of **Example-22E** (200 mg; 1.12 mmol) was added in one portion to this solution. The mixture was stirred until the title product of **Example-22E** 5 was consumed (1-2 hours). The mixture was chilled with an ice-bath and then treated with 6 M HCl (830 L). ¹HNMR analysis indicated a chemical yield of 95 mole% or better. The solvent was evaporated and the title product of **Example-22** was purified by reverse-phase or ion-exchange chromatography.

10 A 210gm solution (containing ~20 g of the title product of **Example-22E** of the base hydrolysis reaction product was put into a 500 mL, three-necked, round-bottomed flask. The apparatus was equipped with a mechanical stirrer, a Dean-Stark apparatus (20 mL with a stopcock), a condenser, and a temperature controller. Water (140 mL) was distilled off from the mixture. 1-butanol (150 mL) was added to the pot and the remaining water (37 mL) was 15 distilled azeotropically. Additional 1-butanol (13 mL) was removed by distillation until the pot temperature reached 117 °C. The butanol slurry was cooled to room temperature and filtered through a pad of celite. The salts were washed with 1-butanol (2x20 mL). DBU (21.8 L; 146 mmol) and ethyl acetimidate hydrochloride (17.1 mg; 134 mmol) were dissolved in 1-butanol (40 mL) in a 500 mL, three-necked, round-bottomed flask at room 20 temperature. The apparatus was equipped with a mechanical stirrer, an addition funnel, and a temperature probe. The title product of **Example-22E** /1-butanol solution was put into the addition funnel and added to the ethyl acetimidate / DBU solution while maintaining the pot temperature below 25 °C. The mixture was stirred until the starting material was consumed (2-3 hours). A solution of conc. HCl (94 mL) and water (100 mL) was put into a 1 L, three- 25 necked, round-bottomed flask and chilled to 0 °C. The apparatus was equipped with a mechanical stirrer, an addition funnel, and a temperature probe. The reaction mixture was put into the addition funnel. The reaction mixture was added to the aqueous HCl solution while maintaining the temperature below 25 °C. Ethyl acetate (100 mL) was added to the solution and the layers were separated. The aqueous layer was washed once more with ethyl 30 acetate (100mL). ¹HNMR analysis indicated a chemical yield of 95 mole% or better. This title product of **Example-22** was purified by reverse-phase or ion-exchange chromatography.

¹HNMR (400MHz, D₂O) 1.49 (3H, s), 2.08 (3H, s), 2.74 (2H, m), 2.91 (1H, d), 3.17 (1H, d), 3.35 (2H, t).

Example 23

5



S-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine methyl ester, dihydrochloride

10 The product of **Example 22** (1.0 g, 3.42 mmol) dissolved in anhydrous methanol (40ml) was added to a 500 ml 3 neck round bottom flask equipped with a magnetic stirrer and a thermocouple. This reaction under nitrogen was cooled to 0 °C. HCl gas was bubbled into the reaction solution for 1 minute. The reaction mixture was allowed to warm to room temperature and continued to stir overnight. A sample was taken from the reaction mixture and concentrated. NMR and Mass Spectrometry indicated starting material and product.

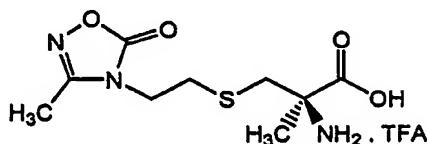
15 The solvent was stripped and the oily residue was re-dissolved in anhydrous methanol (40 ml), cooled to 0 °C, and HCl gas was bubbled into the solution for 1 minute. The reaction mixture was allowed to warm to room temperature and stir overnight. A sample was taken from the reaction mixture and concentrated. NMR and Mass Spectrometry indicated minimal starting material and majority product. The solvent was stripped and the oily residue was re-dissolved in anhydrous methanol (40 ml), cooled to 0 °C, and HCl gas was again bubbled into the solution for 1 minute. The reaction mixture was allowed to warm to room temperature and stir overnight. A sample was taken from the reaction mixture and concentrated. NMR and Mass Spectrometry indicated only the desired title product. The reaction mixture was concentrated to provide 1.01 g of a light yellow oil, 97% yield. The reaction mixture was stirred in acetonitrile (50 ml) for 3 hours and the title product as a white fine powder was recovered, 484 mg. Mass Spectrometry: (ZMD Waters Micromass, Electrospray), M+H at 234.2

¹HNMR: (400mHz, D₂O) δ 1.51 (s, 3H), 2.09 (s, 3H), 2.72 (t, 2H), 2.97 (d, 1H), 3.19 (d, 1H), 3.36 (t, 2H), 3.73 (s, 3H). ¹³CNMR: δ 18.58, 21.69, 30.79, 37.79, 41.58, 54.24, 60.75, 165.41, 171.35.

Analysis calculated for C₉H₁₉N₃O₂S + 2 HCl + 0.3 H₂O (311.66): C 34.68, H 6.98, N 13.48,

5 Cl 22.75, S 10.29. Found: C 34.51, H 6.99, N 13.75, Cl 22.75, S 10.43.

Example 24

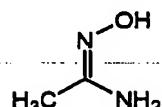


10

**2-methyl-S-[2-(3-methyl-5-oxo-1,2,4-oxadiazol-4(5H)-yl)ethyl]- L-cysteine,
monotrifluoroacetate**

Example-24A) *N*¹-hydroxyethanimidamide

15



To a 3L round bottom flask was added hydroxylamine hydrochloride (138.98 g, 2.0 mol)

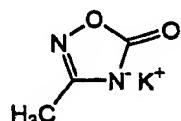
20 in ethanol (1.2L) followed by a slow addition of sodium ethoxide (136.1 g, 2.0mol). The temperature was maintained between 25 °C and 30 °C. The reaction was then stirred for 30 minutes at room temperature. The precipitated NaCl was filtered off and washed with ethanol (100 mL). The hydroxylamine free base in the filtrate was added to a 3L flask and acetonitrile (112.75 g, 2.75 mol). This mixture was then subjected to reflux over night.

25 After cooling, the solvent was carefully removed in vacuo to 50% of its original volume. The reaction then sat in an ice bath for one hour were upon crystals formed and were filtered off. The filtrate was carefully concentrated to 50% of its volume again. The reaction was

placed into ice and the resulting crystals were again isolated by filtration to afford 52 g (35%) of the title product.

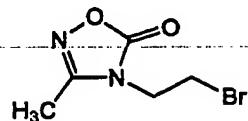
Example-24B) potassium 3-methyl-1,2,4-oxadiazolin-5-onate

5



To a 25 mL round bottom flask was added the product of **Example-24A** (1 g, 0.013 mol), potassium t-butoxide (1.59 g, 0.013 mol) and diethyl carbonate (8.18 mL, 0.067 mol). The 10 reaction was then refluxed for 5 hours. The solvent was removed and the resulting solid was triturated with methylene chloride and diethyl ether. The solid title product was then dried under high vacuum to afford 1.57 g (87%). ^1H NMR (d_6 -DMSO, 300MHz) δ 1.69 (bs, 3H). ^{13}C NMR (d_6 -DMSO, 99MHz) δ 13.26, 166.54, 173.99.

15 **Example-24C) 3-methyl-1-(1-bromoethyl)-2,4-oxadiazolin-5-one**

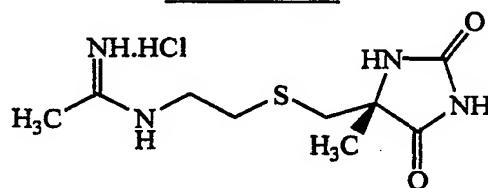


To a 250 mL Round Bottom Flask was added the title product of **Example-24B**, 20 potassium 3-methyl-1,2,4-oxadiazolin-5-onate, (10.13 g, 0.0734 mol) in DMF (100mL). To the slurry was added 1,2-dibromo ethane (31.54 mL, 0.366 mol). The reaction was heated in an oil bath for 2 hours at 130 °C. The oil bath was removed and the reaction cooled after which water (200 mL) and ethyl acetate (50mL) were added. The organics were collected and washed 3X100 mL brine. The organics were dried over MgSO_4 and then concentrated 25 in vacuo to afford 9.1 g (60%) of the title compound.

^1H NMR (CDCl_3 , 300MHz) δ 2.21 (s, 3H), 3.12 (t, 2H), 3.91 (t, 2H).

The 75mL of methanol in a 100 mL round bottom flask was deoxygenated by bubbling nitrogen through it for 5 minutes. To 50 mL of this methanol was added NaOH (1.6 g, 0.040 mol). The suspension was stirred in an oil bath at 45 °C for 30 minutes after which the NaOH had dissolved. The resulting solution was cooled to room temperature and alpha-5 methyl cysteine (1.72 g, 0.010 mol) was added in 10 mL of deoxygenated methanol. The reaction stirred for 45 minutes at room temperature. To this reaction was added the product of Example-24C (2.07 g, 0.010 mol) in 10 mL of deoxygenated methanol. The reaction was complete as indicated by mass spectral analysis after it had been stirred over night. The reaction mixture was diluted with water (100 mL) and purified using reverse phase 10 chromatography to afford 3.0 g (93%) of the title product of Example 24 as its trifluoroacetate salt. M.S. M+H⁺(262.0), M+Na⁺(282.0). ¹H NMR (CD₃OD, 300MHz) δ 1.39 (s, 3H), 2.23 (s, 3H), 2.74 (m, 2H), 2.84 (m, 2H), 3.72 (t, 2H).

15

Example 25

[2-[[[(4R)-4-methyl-2,5-dioxo-4-imidazolidinyl]methyl]thio]ethyl](1-NH₂)iminoethyl)amine

20 The [2-[[[(4R)-4-methyl-2,5-dioxo-4-imidazolidinyl]methyl]thio]ethyl]carbamic acid, 1,1-dimethylethyl ester isomer product of Example-22D (2.05 g, 6.5 mmol) was dissolved in 25mL 4.0 N HCl in Dioxane and stirred for 10 min. After the addition of 2N HCl (5 mL) the reaction was stirred for additional 2 h. The reaction mixture was then concentrated under reduced pressure to give 1.68 g of a red brown gummy solid. This material was taken up in 25 25 mL of deionized water and the pH was adjusted to 8.4 with 2N NaOH. Ethylacetimidate hydrochloride (2.39 g, 0.019 mol) was then added while maintaining the pH at 8.4. The reaction mixture was then stirred at room temperature for one hour at pH 8.4. The pH of the reaction mixture was then adjusted to 3.5 by adding an appropriate amount of 1 N HCl and stirred for another 16 h. The reaction mixture was then concentrated on a rotary evaporator

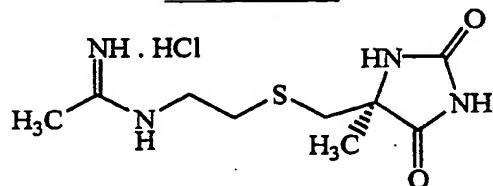
to obtain the crude product that was purified on a Gilson preparative HPLC to give the desired product as a white hygroscopic solid in 70% yield.

Mass $M^{+1} = 245$. $[\alpha]$ in H_2O at $25^\circ C = -37.6$ (365 nm).

Analysis calculated for $C_9H_{16}N_4O_2S + 1.0 HCl + 1.3H_2O$ (formula weight = 304.20): C

5 35.54, H 6.49, N 18.42, Cl 11.65, S 10.54. Found: C 35.83, H 6.08, N 18.55, Cl 11.19, S 10.63.

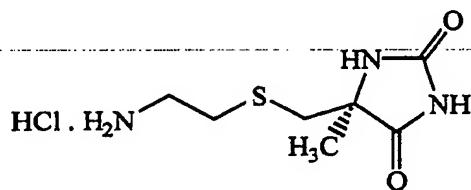
Example 26



10

[2-[[[(4S)-4-methyl-2,5-dioxo-4-imidazolidinyl]methyl]thio]ethyl](1-N-iminoethyl)amine hydrochloride

15 **Example-26A) (5S)-5-[[((2-aminoethyl)thio)methyl]-5-methyl-2,4-imidazolidinedione, monohydrochloride**



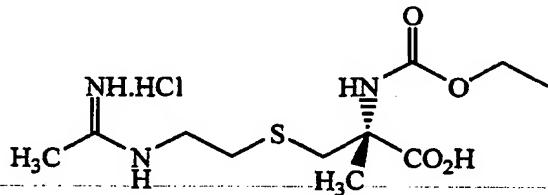
20 The [2-[[[(4S)-4-methyl-2,5-dioxo-4-imidazolidinyl]methyl]thio]ethyl]carbamic acid, 1,1-dimethylethyl ester isomer product of Example 22D was purified by chromatography using 66 % Ethyl Acetate in Toluene, Biotage Flash 75 silica gel. A sample of this material (5.9 g, 16.5 mmol, $[\alpha]$ in $MeOH$ at $25^\circ C = +45.7$, 365 nm) was then dissolved in 165 mL THF and treated with 4.125 mL 4.0 N HCl in Dioxane. The reaction was allowed to stir for two hours at room temperature and monitored by TLC. The free amine product was then purified 25 by chromatography using reverse phase media (YMC-ODS-AQ) to yield 4.8 g of the title material.

A 3.5 g (17.2 mmol) sample of the product of Example-26A was treated with 10 % NaOH solution to pH 9-10. To this solution was added 4.26 g of ethyl acetimidate hydrochloride while adjusting pH to 9 by adding a solution of 10% NaOH. After stirring at 5 pH 9 for 2 hours, the pH was adjusted to 7.5 by adding an appropriate amount of 0.1 N HCl. This solution was stirred for another 2 hours before the pH was further adjusted to 4.5 by adding 0.1 N HCl. After stirring this solution for 10 hours the water was removed under reduced pressure (11 mbar) and 47°C water bath. The crude title product was chromatographed using reverse phase media (YMC-ODS-AQ) to give 156 mg of the title 10 material. $[\alpha]$ in H_2O at 25 °C = +54.8 (365 nm).

Analysis calculated for $\text{C}_9\text{H}_{16}\text{N}_4\text{O}_2\text{S} + 1.0 \text{ HCl} + 0.85 \text{ H}_2\text{O}$ (formula weight = 296.09): C 36.51, H 6.37, N 18.92, Cl 11.97, S 10.83. Found: C 36.69, H 6.32, N 18.85, Cl 11.46, S 11.12.

15

Example 27

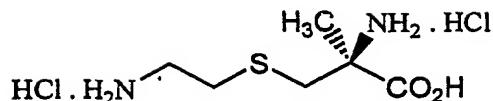


N-(ethoxycarbonyl)-*S*-[2-[(1-iminoethyl)amino]ethyl]-2-methyl-L-cysteine,
monohydrochloride

20

A sample of the product of Example 1 (3.22 g, 0.01 mol) was taken up in 50mL of deionized water and to this K_2CO_3 (2.76 g) was added followed by the addition of ethylchloroformate (1.08 g, 0.01 mol). The reaction mixture was stirred at 25 °C for 1 hour 25 and then concentrated on rotary evaporator to give a white solid. This solid was purified by HPLC to give the desired product.

Mass M^{+1} = 292

Example 28

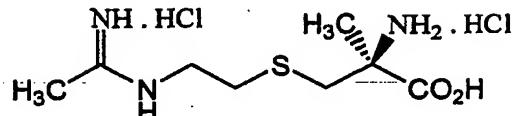
5

S-(2-aminoethyl)-2-methyl-D-cysteine dihydrochloride

A sample of the [2-[[[(4S)-4-methyl-2,5-dioxo-4-imidazolidinyl]methyl]thio]ethyl]carbamic acid, 1,1-dimethylethyl ester product of Example 22D (1.025 g, 3.25 mmol) was dissolved in 10 35mL conc. HCl and stirred for 46 h at reflux temperature. The reaction mixture was then concentrated under reduced pressure to give 900 mg of a red brown gummy solid. This crude product was purified by reverse phase HPLC to give pure S-(2-aminoethyl)-2-methyl-D-cysteine dihydrochloride (800 mg, 98% yield). Mass M⁺ = 179. [α] in H₂O at 25 °C = - 85.6 (365 nm).

15 Analysis calculated for C₆H₁₄N₂O₂S + 2 HCl + 1 H₂O + 1.6 NH₄Cl (formula weight 356.39; exact mass 178.07): C 20.22, H 7.35, N 14.15, Cl 35.81, S 9.00. Found: C 20.09, H 6.95, N 14.55, Cl 36.15, S 9.56.

20

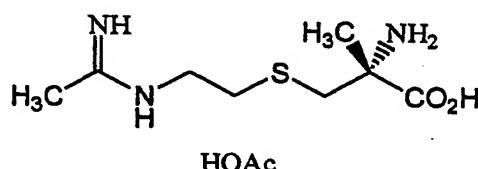
Example 29**[2-(1-Iminoethylamino)ethyl]-2-methyl-D-cysteine hydrochloride**

25 A sample of the product of Example 28 (1.25 g, 0.005 mol) was taken up in 20 mL of de-ionized water and the pH was adjusted to between 8.5 and 9 with 0.1N NaOH. Ethylacetimidate hydrochloride (2.39 g, 0.019 mol) was then added to the stirred reaction while maintaining the pH at 8.5. The reaction mixture was then stirred at 25 °C and 8.5 pH

for 2 hours. The pH of the reaction mixture was then adjusted to 4.0 by adding an appropriate amount of 0.1 N HCl. The reaction mixture was then concentrated on a rotary evaporator and the crude product residue was purified on a Gilson HPLC system using YMC AQ column with 0.1% AcOH/CH₃CN/H₂O to give the desired product in quantitative yield.

5 Mass M⁺= 220. [α] in H₂O at 25 °C = -134.5 (365 nm).
 Analysis calculated for C₈H₁₇N₃O₂S + 1.2 HCl + 2 H₂O (formula weight 299.09; exact mass 219.10): C 32.13, H 7.48, N 14.05, Cl 14.22, S 10.72. Found: C 32.39, H 7.26, N 14.05, Cl 14.33, S 10.42.

10

Example 30**S-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine, acetate**

15

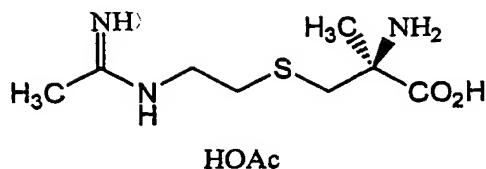
Bio-Rad AG 1x8 resin, 200-400 mesh in the acetate form (300 g, 960 meqv.) was slurried in HPLC grade water and loaded onto a column 8cm in diameter. The water was drained to the top of the column before 37 g (116 mmol) of the product of Example 1, dissolved in 10 mL of water, was loaded onto the column. The material was then eluted with 1 L of water.

20 The first 200 mL fraction contained no product but the subsequent 500 mL yielded 30 g of the desired title product as a white glassy solid after removal of the water under reduced pressure.

Analysis calculated for C₈H₁₇N₃O₂S + CH₃COOH + 1.3 H₂O: C 39.67, H 7.86, N 13.88.
 Found: C 39.96, H 7.87, N 13.69.

25

Example 31

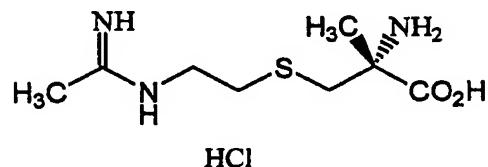


***S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-D-cysteine, acetate**

5 A sample of the product of **Example 29** as its mono hydrochloride salt (101 mg, 0.33 mmol) was converted to the title mono acetate by the method of **Example 30**.
 Analysis calculated for $C_8H_{17}N_3O_2S \cdot CH_3COOH + 0.05 HCl + 2.2 H_2O$: C 37.41, H 8.01, N 13.2, Cl 0.56. Found: C 37.30, H 7.92, N 13.17, Cl 0.41.

10

Example 32



***S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine, monohydrochloride**

15

This material was prepared by passing the product of **Example 1** through a reverse phase column using the conditions described in **Example 28**.
 Analysis calculated for $C_8H_{17}N_3O_2S + 1.05 HCl + 0.8 H_2O$: C 35.35, H 7.36, N 15.44, Cl 13.65. Found: C 35.33, H 7.28, N 15.45, Cl 13.68.

20

Example 33

D-galacturonic acid salt of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine

To a stirred 10 mL 0.001 M solution of acetate salt product of **Example 30** was added D-galacturonic acid monohydrate (0.21 g, 0.001 mole). After stirring for 2 hours, the solution was concentrated under vacuum. The title galacturonic acid salt was dissolved in 10 mL of

water and lyophilized.

Example 34

Succinic acid salt of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-D-cysteine

5

The title material was prepared by the method of **Example 33** from succinic acid and the product of **Example 31**.

Analysis calculated for $C_8H_{17}N_3O_2S + C_4H_6O_4 + 1.5 H_2O$: C 39.55, H 7.19, N 11.53. Found: C 39.24, H 6.04, N 11.41.

10

Example 35

Succinic acid salt of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine

The title material was prepared by the method of **Example 33** from succinic acid and the product of **Example 30**.

Analysis calculated for $C_8H_{17}N_3O_2S + C_4H_6O_4 + 1.1 H_2O$: C 39.99, H 7.40, N 12.33. Found: C 40.35, H 7.11, N 11.76.

20

Example 36

Ethanolamine salt of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine

The title material was prepared by the method of **Example 33** from ethanolamine and the product of **Example 30**.

Analysis calculated for $C_8H_{17}N_3O_2S + C_2H_7NO + 2HCl + 1.3 H_2O$: C 31.73, H 7.67, N 14.80, Cl 18.73. Found: C 31.41, H 7.60, N 15.00, Cl 19.12.

Example 37

Ethylene diamine salt of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine

30 The title material was prepared by the method of **Example 33** from ethylene diamine and the product of **Example 30**.

Analysis calculated for $C_8H_{17}N_3O_2S + 2HCl + C_2H_8N_2 + 1.2 H_2O$: C 32.12, H 7.92, N 18.73, Cl 18.96. Found: C 31.90, H 9.19, N 18.08, Cl 19.11.

Example 38

5 **DL-Aspartic acid salt of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine**

The title material was prepared by the method of **Example 33** from DL-aspartic acid and the product of **Example 30**.

Analysis calculated for $C_{12}H_{24}N_4O_6S + 1.8 H_2O + 0.4 HOAc$ (formula weight = 408.86):

10 C 37.60, H 7.20, N 13.70. Found: C 37.59, H 7.66, N 13.73.

Example 39

D-Glutamic acid salt of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine

15 The title material was prepared by the method of **Example 33** from D-glutamic acid and the product of **Example 30**.

Analysis calculated for $C_{13}H_{26}N_4O_6S + 1.8 H_2O + 0.3 HOAc$ (formula weight = 416.88): C 39.18, H 7.45, N 13.44. Found: C 39.47, H 7.52, N 13.29.

20

Example 40

Citric acid salt of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine

The title material was prepared by the method of **Example 33** from citric acid and the product of **Example 30**.

25 Analysis calculated for $C_{14}H_{25}N_3O_9S + 0.5 H_2O + 0.1 HOAc + 0.15 EtOH$ (formula weight = 433.36): C 40.19, H 6.35, N 9.70. Found: C 40.32, H 5.74, N 9.58.

Example 41

DOWEX 50WX4-400 ion-exchange resin salt of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine

30

DOWEX® 50WX4-400 (3 g, 1.6 meq/mL, 4.8 meq/g) was washed with deionized water (all water used in this experiment was deionized) to pH 6 of the wash. The resin was dried for one hour at room temperature. The product of **Example 30** (0.6 g) in 30 mL of water was added to DOWEX resin (0.2 g). The suspension was shaken for three hours at ambient

5 temperature using an ORBIT™ shaker, then stripped to dryness. This procedure was repeated three times with fresh 30 mL of water added after each concentration of the reaction mixture. With the final portion of fresh water, the slurry was shaken overnight at ambient temperature

After stripping the reaction to dryness, 15 mL of water were added. The resin was filtered and washed three times with additional 15 mL of water. The filtrate was concentrated

10 (several drops of acetic acid were added) and dried under vacuum giving 0.4 g of starting SC-84250 which was confirmed by ¹H NMR (D₂O). The loaded resin was dried at ambient temperature on the bench, followed by 1 hour under vacuum giving 0.3 g of loaded resin. A sample of this resin and a sample of unreacted washed DOWEX 50WX4-400 were

15 submitted for nitrogen combustion analysis: The results for the untreated resin were % of N is 0 %; for the loaded resin % of N is 9.72 %.

Example 42

Potassium hydrogen sulfate acid salt of S-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine

20

The title material was prepared by the method of **Example 33** from 0.001 mole of KHSO₄ and the product of **Example 30**.

Analysis calculated for C₈H₁₇N₃O₂S + KHSO₄ + 2 H₂O: C 24.75, H 5.68, N 10.90, S 16.00.

Found: C 24.54, H 5.66, N 10.73, S 16.38.

25

Example 43

Potassium hydrogen sulfate acid salt of S-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine

30

The title material was prepared by the method of **Example 33** from 0.001 mole of KHSO₄ and the product of **Example 30**.

Example 44**Hydrogen sulfate acid salt of S-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine**

To a stirred 10 mL solution of the product of **Example 30** was added 2 mL 0.505 N H₂SO₄. After stirring for 2 hours, the solution was concentrated under vacuum. The resulting salt was dissolved in 10 mL of water and lyophilized. Analysis calculated for C₈H₁₇N₃O₂S + 0.5 H₂SO₄ + 1.5 H₂O: C 32.58, H 7.23, N 14.75, S 16.42. Found: C 32.53, H 7.17, N 14.23, S 16.28.

10

Example 45**Glycerate salt of S-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine**

S-Glyceric acid was prepared from its calcium salt by stirring for 4 hours with Dowex® 50W resin in its acid form. The resin was filtered and washed with H₂O. The resulting filtrate was concentrated and dried under vacuum. Analysis calculated for C₃H₆O₄: C 31.31, H 6.13. Found: C 31.29, H 6.19. The method described in **Example 33** was used to prepare the S-glyceric acid salt from the product of **Example 30** starting with 0.001 mole of S-glyceric acid. Analysis calculated for C₁₁H₂₃N₃O₆S + 1.5 H₂O: C 37.49, H 7.44, N 11.92, S 9.10. Found: C 37.49, H 7.31, N 11.73, S 9.22.

25

Example 46**Malate salt of S-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine**

The method described in **Example 33** was used to prepare the malic acid salt from the product of **Example 30** starting with 0.001 mole of malic acid. Analysis calculated for C₈H₁₇N₃O₂S + 1.33 H₂O + C₄H₆O₅: C 38.20, H 6.85, N 11.15. Found: C 38.37, H 6.51, N 11.09.

30

Example 47**Hemi-malate salt of S-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine**

The method described in **Example 33** was used to prepare the malic acid salt from the product of **Example 30** starting with 0.0005 mole of malic acid.

Analysis calculated for $C_8H_{17}N_3O_2S + 1.75 H_2O + 0.5 C_4H_6O_5$: C 37.92, H 7.48, N 13.22.

Found: C 37.92, H 7.88, N 13.03.

5

Example 48

Potassium dihydrogen phosphate salt of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine

10 The method described in **Example 33** was used to prepare the title salt from the product of **Example 30**.

Analysis calculated for $C_8H_{17}N_3O_2S + KH_2PO_4 + 2.5 H_2O + 0.66 HOAc$: C 25.44, H 6.10, N 9.55. Found: C 25.27, H 5.95, N 9.80.

15

Example 49

Sodium dihydrogen phosphate salt of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine

20 The method described in **Example 33** was used to prepare the title salt from the product of **Example 30**.

Analysis calculated for $C_8H_{17}N_3O_2S + NaH_2PO_4 + 2 H_2O + 0.3 HOAc$: C 26.26, H 6.20, N 10.68. Found: C 26.57, H 6.25, N 10.72.

25

Example 50

Calcium dihydrogen phosphate salt of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine

The method described in **Example 33** was used to prepare the title salt from the product of **Example 30**.

30 Analysis calculated for $C_8H_{17}N_3O_2S + 2 NaH_2PO_4 + 2 H_2O$: C 19.40, H 5.09, N 8.48. Found: C 19.34, H 5.10, N 8.56.

Example 51**Calcium phosphate salt of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine**

5 The method described in **Example 33** was used to prepare the title salt from the product of **Example 30**.

Analysis calculated for $C_8H_{17}N_3O_2S + Ca(H_2PO_4)_2 + 0.2 HOAc$: C 19.96, H 4.35, N 8.31.

Found: C 20.14, H 5.73, N 8.80.

Example 52**10 Calcium hydrogen phosphate salt of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine**

The method described in **Example 33** was used to prepare the title salt from the product of **Example 30**.

15 Analysis calculated for $C_8H_{17}N_3O_2S + CaHPO_4 + 2.2 HCl + H_2O$: C 21.18, H 4.93, N 9.26.

Found: C 21.20, H 5.28, N 9.37.

Example 53**20 Calcium phosphate, tribasic salt of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine (1.62:1)**

The method described in **Example 33** was used to prepare the title salt from the product of **Example 30**.

Analysis calculated for $C_8H_{17}N_3O_2S + Ca_3(PO_4)_2 + HOAc + 3 H_2O$: C 14.37, H 2.77, N 5.03.

25 Found: C 14.13, H 3.01, N 4.71.

Example 54**Calcium phosphate, tribasic salt of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine (1:1)**

30

The method described in **Example 33** was used to prepare the title salt from the product of **Example 30**.

Analysis calculated for $C_8H_{17}N_3O_2S \cdot Ca_3(PO_4)_2 + 2.2 HCl + 2H_2O$: C 14.88, H 3.62, N 6.51.
 Found: C 15.09, H 3.85, N 6.23.

Example 55

5 Bio-Rex® 70 salt of S-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine

The method described in **Example 41** was used to prepare the Bio-Rex® 70 salt of the neutral form of the product of **Example 1**. The results for the untreated resin were % of N is 0 %; for the loaded resin % of N is 7.93 %.

10

Example 56

IPR (amberlite)-69 salt of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine

The IPR-69 salt of the neutral form of the product of **Example 1** was prepared in the same manner as described in **Example 41** except the resin was treated first with 1 N HCl to convert it to H⁺ form. This resin is a polystyrenedivinylbenzene sulfonic acid resin as is the Dowex-50. However, it is GMP quality but covers a broader mesh size. It is less colored than the Dowex. After the washings and prior to loading the compound, the resin was slurried in H₂O and the fine particles that rose to the top were decanted. From this reaction was recovered 4.9 g of salt. 7.69 % N (or 0.401 g of SC-84250/g of resin).

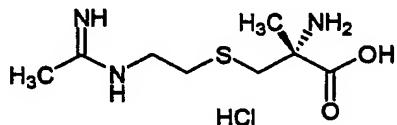
Example 57

IPR (amberlite)-69 salt of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine

25 The IPR-64 salt the neutral form of the product of **Example 1** was prepared in the same manner as described in **Example 41** with the decanting of fines as described in **Example 56**. This resin is the same as the Bio-Rex 70 with the exception that is GMP quality. The only variation was after shaking overnight the resin was shaken an additional two times with strips in between. From this reaction was recovered 4.3 g. 6.20 % N (0.346 g of
30 compound/g of resin

Example 58

**Preparation of monohydrochloride from the dihydrochloride.
salt of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine**



5

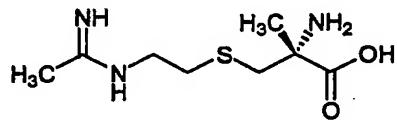
Dissolve ~120 mg of the product of **Example 1** in 3 ml of DMF. Add 1 ml of propylene oxide and stir. The product will precipitate. Wash with ether. Dissolve product in water and freeze dry. Table 1 shows the elemental analysis.

10

Example 59

Preparation of monosubstituted salts of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine salts by AgCl precipitation

15



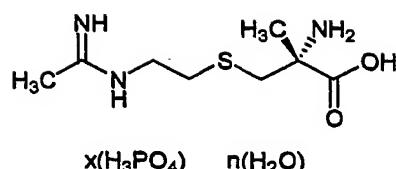
R n (H₂O)

Dissolve the product of **Example 58** in water. Add a stoichiometric amount of a silver salt. Filter the solids. Freeze dry the remaining solution. Table 2 shows the elemental analyses. 20 where n represents the moles of water.

Example 60

Preparation of phosphate salts of *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine salts by AgCl precipitation

25

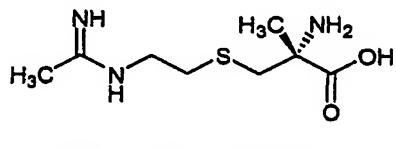


Dissolve the product of **Example 1** in water. Add two moles of Ag_3PO_4 per mole of product of **Example 1** and mix. Filter out the solids and freeze dry the resulting solution. Analyze 5 the resulting material, and adjust the phosphate content with H_3PO_4 . Table 3 shows the elemental analyses where x represents the moles of phosphoric acid.

Example 61

10

Preparation of mixed salts from *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine monohydrochloride



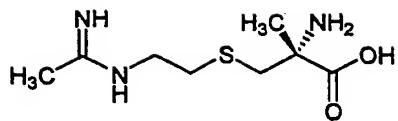
15

Dissolve the product of **Example 58** in water. Add a stoichiometric amount of reagent (R). Table 4 shows the elemental analyses where x represents the moles of R.

20

Example 62

Preparation of mixed salts from *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine dihydrochloride

2HCl nH₂O

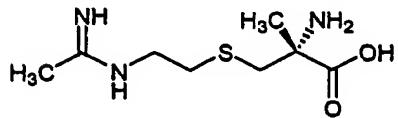
Dissolve the product of **Example 1** in water. Add base until the pH is 6. Table 5 shows the elemental analyses.

5

Example 63

Preparation of zinc salts from S-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine dihydrochloride

10

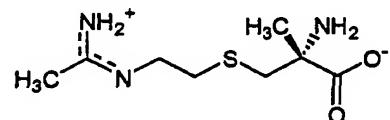


(ZnO) 2 (HCl))

Dissolve the product of **Example 1** in water. Mix with excess zinc oxide. Filter and freeze
15 dry the resulting solution. Table 6 shows the elemental analysis.

Example 64

20 Preparation of mixed salts from S-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine neutral compound

x(MA) n(H₂O)

Dissolve the neutral form of the product of **Example 20** in water. Mix with desired reagent and freeze dry. Table 7 shows the elemental analyses where x represents the moles of MA, metal cation and counter anion.

5

Example 65

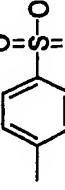
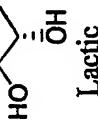
Preparation of salts from *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine neutral compound

The following Table 8 lists salts and their elemental analyses prepared from the product of **Example 1** by one of the variety of methods described in this application. Table 8 illustrates the elemental analysis of these salts where D represents *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine with the formula of $C_8H_{17}N_3O_2S$ and A the empirical formula of the indicated acid and or counter ion source.

Table 1.

Formula		Calculated	Found				
	C	H	N	Cl	C	H	Cl
$\text{C}_8\text{H}_{17}\text{N}_3\text{O}_2\text{S} (\text{HCl}) (\text{H}_2\text{O})$	35.10	7.36	15.35	12.95	35.06	7.53	14.90
							13.07

Table 2.

Acid (R)	Formula	Calculated	Found				
		C	H	N	S	C	
$\text{H}_3\text{C}-\overset{\text{O}}{\underset{\text{O}}{\text{S}}}-\text{OH}$	$\text{C}_8\text{H}_{17}\text{N}_3\text{O}_2\text{S}$ (CH_3SO_3) (H_2O)	32.42	6.95	12.60	19.23	32.09	7.06
Methane Sulfonic							12.36
							19.67
	$\text{C}_8\text{H}_{17}\text{N}_3\text{O}_2\text{S}$ $\text{C}_7\text{H}_8\text{O}_3\text{S}$ (H_2O)	43.99	6.65	10.26	15.66	43.83	6.75
Toluene Sulfonic							9.91
							14.5
	$\text{C}_8\text{H}_{17}\text{N}_3\text{O}_2\text{S}$ $\text{C}_3\text{H}_6\text{O}_3$ (H_2O)	40.36	7.70	12.83	9.79	40.79	7.84
Lactic							12.6
							9.68

HNO ₃ Nitric	C ₈ H ₁₇ N ₃ O ₂ S HNO ₃ (H ₂ O)	31.99	6.71	18.66	10.67	32.32	5.79	18.32	-
Acetic	C ₈ H ₁₇ N ₃ O ₂ S C ₂ H ₄ O ₂ 1.5(H ₂ O)	39.20	7.90	13.72	10.46	39.19	7.15	13.53	-
Benzoic	C ₈ H ₁₇ N ₃ O ₂ S C ₇ H ₆ O ₂ 1.5(H ₂ O)	48.90	7.11	11.40	8.70	48.93	7.45	11.74	-
Pamoic	C ₈ H ₁₇ N ₃ O ₂ S .5(C ₂₃ H ₁₆ O ₆) 1.5(H ₂ O)	53.17	6.41	9.54	7.28	53.65	6.38	9.92	6.92

Table 3.

$\text{C}_8\text{H}_{17}\text{N}_3\text{O}_2\text{S} (\text{H}_3\text{PO}_4) 1.5(\text{H}_2\text{O})$	27.91	6.73	12.20	9.00	27.74	6.03	12.12	8.73
$\text{C}_8\text{H}_{17}\text{N}_3\text{O}_2\text{S} 2(\text{H}_3\text{PO}_4) 2(\text{H}_2\text{O})$	21.29	6.03	9.31	13.73	20.89	5.64	9.3	13.67

Table 4.

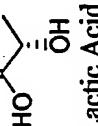
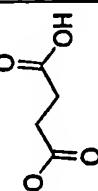
Formula	Calculated				Found			
	C	H	N	Cl	C	H	N	Cl
CaCl_2	28.41	6.26	12.42	20.96	28.42	6.3	12.26	19.6
$\text{C}_8\text{H}_{17}\text{N}_3\text{O}_2\text{S}$.5(CaCl_2) 1.5(H_2O) (HCl)								
$\text{C}_8\text{H}_{17}\text{N}_3\text{O}_2\text{S}$ $\text{C}_3\text{H}_6\text{O}_3$ HCl	38.2	6.99	12.15	10.25	38.32	7.16	12.23	10.81
 Lactic Acid								
$\text{C}_8\text{H}_{17}\text{N}_3\text{O}_2\text{S}$.5($\text{C}_4\text{H}_6\text{O}_4$) HCl .5(H_2O)	37.09	6.85	12.98	10.95	37.22	6.68	13.08	11.45
 Succinic Acid								

Table 5.

Formula	Calculated					Found				
	C	H	N	M	Cl	C	H	N	M	Cl
C ₈ H ₁₆ N ₃ O ₂ S Li 2(HCl) 2(H ₂ O)	28.75	6.64	12.57	2.08	21.22	28.05	6.58	12.34	2.14	21.59
C ₈ H ₁₆ N ₃ O ₂ S Na 2(HCl) 1.5(H ₂ O)	28.16	6.20	12.31	—	20.78	27.71	6.10	12.47	—	22.20
C ₈ H ₁₆ N ₃ O ₂ S K 2(HCl) 2(H ₂ O)	26.23	6.05	11.47	—	19.36	25.75	5.60	11.50	—	21.13

Table 6.

Formula/ References	Calculated					Found				
	C	H	N	Cl	Zn	C	H	N	Cl	Zn
C ₈ H ₁₇ N ₃ O ₂ S 2(HCl) ZnO	25.72	5.13	11.25	18.98	17.50	25.98	4.52	11.57	20.54	17.13

Table 7.

Formula	Measured			Found		
	C	H	N	C	H	N
$C_8H_{17}N_3O_2S$ (NaCl) 2(H ₂ O)	30.62	6.75	13.39	30.93	6.39	13.27
$C_8H_{17}N_3O_2S$ 0.5(CaCl ₂) 1.5(H ₂ O)	31.84	6.68	13.92	31.59	6.57	13.73
$C_8H_{17}N_3O_2S$ (NaCH ₃ SO ₃) 2(H ₂ O)	28.95	6.48	11.25	28.91	6.06	10.99

Table 8.

Acid / Formula	Calculated			Found		
	C	H	N	C	H	N
L-Tartaric Acid						
D * 0.6A * 0.75 H ₂ O	38.59	6.91	12.98	38.67	6.94	12.65
D * 1.0A * 0.5 H ₂ O	38.09	6.39	11.10	38.04	6.43	11.38
D * 1.2A * 2.0 H ₂ O	32.48	5.75	8.88	32.76	5.50	8.96
D * 1.1A * 1.5 H ₂ O	34.37	5.93	9.70	34.51	5.81	9.69
D-Tartaric Acid						
D * 0.5A * 0.5 H ₂ O	39.59	6.98	13.85	39.27	7.09	13.34
D * 1.0A * 1.0 H ₂ O	37.20	6.50	10.85	37.85	6.45	10.43
(R)-(-)-Mandelic Acid						
D * 1.0A * 2.5 H ₂ O	49.99	7.87	10.93	49.90	6.92	10.83
(S)-(+)-Mandelic Acid						
D * 1.0A * 2.75 H ₂ O	49.41	7.90	10.86	49.38	8.23	10.78

Citric Acid	40.22	7.08	14.23		40.96	6.87	13.26	
D * 0.4A * 0.8 H ₂ O	40.19	6.35	9.70		40.32	5.74	9.58	
D * 1.0A * 0.5 H ₂ O * 0.1 HOAc * .015 EtOH								
Muic Acid				S				S
D * 0.5A * 2.0 H ₂ O	36.66	7.27	11.66		36.46	7.12	11.24	
D * 0.6A * 1.0 H ₂ O	40.73	6.84	12.95	9.89	38.09	7.13	12.03	9.30
Maleic Acid								
D * 0.5A * 1.5 H ₂ O	39.46	7.29	13.81		39.64	6.98	13.01	
Malonic Acid								
D * 0.5A * 1.5 H ₂ O	38.24	7.34	14.08		38.03	7.37	13.89	

Benzoic Acid								
D *1.1A *2.75 H ₂ O	50.81	7.90	11.32		50.54	6.39	11.12	
D *1.0A *1.5 H ₂ O	48.90	7.11	11.40		48.93	7.45	11.74	
D *2.4 A *1.4 H ₂ O	55.41	6.41	7.82		55.75	6.51	7.52	

Methanesulfonic Acid											
D *1.0A *1.25 HCl *0.8 H ₂ O	28.80	6.40	11.19	11.80	17.08	28.29	6.38	11.16	12.01	17.17	
D *2.0A *0.75 HCl *5.0 H ₂ O	23.10	6.74	8.08	5.11	18.50	23.03	6.58	8.16	5.18	18.58	
D *1.0A *1.0 H ₂ O	32.42	6.95	12.60		19.23	32.09	7.06	12.36		19.67	
D *1.0A- *2.0 H ₂ O *1.0 Na ⁺	28.95	6.48	11.25		28.91	6.06	10.99				
D *1.0A *1.0 Me ₄ N ⁺ *1.75 H ₂ O	37.17	8.52	13.34		15.72	37.45	7.92	12.81		14.95	

Tonic Acid								
D *1.0A *1.5 H ₂ O	45.07	6.82	9.35		45.60	6.55	9.21	
D *2.2A *2.4 H ₂ O	46.36	6.55	6.93		46.76	6.01	6.65	
D *1.0A *1.5 HCl *1.0 H ₂ O	38.81	6.19	9.05	11.46	13.18	38.90	6.37	8.93
D *1.0 *1.0 H ₂ O	43.99	6.65	10.26		15.66	43.83	6.75	9.91
D *1.0A- * 1.0 Me ₄ N+ *1.0 H ₂ O	47.28	7.94	11.61		13.29	47.39	7.90	10.40
D *1.0 A *1.0A- * 1.0 Me4N+ *1.0 H ₂ O	47.83	6.79	8.58		14.73	47.95	7.25	8.49
D *1.2 A *1.0 H ₂ O	42.74	6.18	9.84			42.94	5.83	9.34
D *2.1 A *5.0 H ₂ O	38.57	6.22	6.55			38.87	5.75	6.50

Sulfonacetic Acid D *1.1A *1.1 H ₂ O	33.42	5.89	11.69	17.84	31.61	6.04	10.92		17.35
1,3-Propanedisulfonic Acid D *0.67A *1.0 HCl *1.25 H ₂ O	28.96	6.28	10.12	8.54	18.08	28.72	6.32	10.10	8.96
D *0.3A *1.6 HCl *1.25 H ₂ O	29.34	6.56	11.40	15.39	14.53	29.17	6.71	11.50	15.48
L-(+)-Lactic Acid D *1.0A *1.0 H ₂ O	40.36	7.70	12.83		9.79	40.79	7.84	12.60	9.68
D *1.0A *1.0 H ₂ O *1.0 HCl	38.20	6.99	12.15	10.25		38.32	7.16	12.23	10.81
Nitric Acid D *1.0A *1.0 H ₂ O	31.99	6.71	18.66			32.32	5.79	18.32	

Acetic Acid									
D *1.0 A *1.0 H ₂ O	39.20	7.90	13.72		39.19	7.15	13.53		
D *1.0 A *1.3 H ₂ O	39.67	7.86	13.88		39.96	7.87	13.69		
D *1.0 A * 0.05 HCl *2.2 H ₂ O (S-Enantiomer)	37.41	8.01	13.20	0.56	37.30	7.92	13.17	0.41	
Pamoic Acid									
D *0.5 A *1.5 H ₂ O	53.17	6.41	9.54		7.28	53.65	6.38	9.92	6.92

			P		P		P
Phosphoric Acid							
D *1.0 A-*1.5 H ₂ O	27.21	6.73	12.20	9.00	27.74	6.03	12.12
D *2.0A *2.0 H ₂ O	21.29	6.03	9.31	13.73	20.89	5.64	9.30
D *1.0 A-*2.5 H ₂ O *0.66 HOAc *1.0 K ⁺	25.44	6.10	9.55		25.27	5.95	9.80
D *1.0 A-*2.0 H ₂ O *0.33 HOAc *1.0 Na ⁺	26.26	6.20	10.68		26.57	6.25	10.72
D *2.0 A-*0.2H OAc *1.0Ca++	19.96	4.35	8.31		20.14	5.73	8.80
D *1.0 A-*2.2 HCl *1.0 H ₂ O *0.66HOAc *1.0 Ca++	21.18	4.93	9.26		21.20	5.28	9.37
D *2.0 A---*3.0 H ₂ O *1.0 HOAc *2.0 Ca++	14.37	2.77	5.03		14.13	3.01	4.71
D *2.0 A---*2.2 HCl *2.0 H ₂ O *0.66HOAc *3.0 Ca++	14.88	3.62	6.51		15.09	3.85	6.23

Sulfuric Acid	24.75	5.68	10.90	16.00	24.54	5.66	10.73	16.38
D *1.0 A-*2.0 H ₂ O *1.0 K+	32.58	7.23	14.75	16.42	32.53	7.17	14.23	16.28
D *0.5 A *1.5 H ₂ O	25.97	6.97	27.25	13.86	26.12	6.44	27.32	13.31
D *1.0 A--*2.0 CN3H6+*1.5 H ₂ O	30.50	7.39	23.71	13.57	30.94	6.71	23.87	13.23
D *1.5 A--*.0 CN3H6+*0.75 H ₂ O	37.49	7.44	11.92	9.10	37.49	7.31	11.73	9.22
(S)-Glyceric Acid								
D *0.5 A *1.5 H ₂ O	38.20	6.85	11.15		38.37	6.51	11.09	
L-(-)-Malic Acid								
D *1.0 A *1.33 H ₂ O	37.92	7.48	13.22		37.92	7.88	13.03	
D *0.5 A *1.75 H ₂ O								
L-Ascorbic Acid								
D *1.0 A *2.2 H ₂ O	38.65	6.81	9.66		38.91	6.80	9.43	

3-(N-morpholino)propane Sulfonate D *1.0 A *2.5 H ₂ O	38.04	7.87	11.83	38.41	8.12	11.64
L-cysteic Acid D *0.5 A *2.3 H ₂ O	30.73	6.71	13.03	30.91	7.05	12.98
(4S)-hydroxy-L-proline D *1.0 A *0.3 H ₂ O	44.89	7.53	15.74	44.29	7.60	15.91
Cyclopropane-1,1'- dicarboxylic Acid D *1.0 A *1.3 H ₂ O	41.88	6.92	11.27	42.02	6.68	10.83
D *0.5 A *1.5 H ₂ O	40.50	7.45	13.48	40.70	7.10	12.98
2,2-dimethylmalonic Acid D *1.0 A *2.2 H ₂ O	39.93	7.58	10.75	39.88	7.37	10.37
D *0.5 A *1.8 H ₂ O	39.68	7.80	13.22	40.10	7.81	13.38

Ethanolamine D *0.5 A *1.3 H ₂ O * 2.0HCl	31.73	7.67	14.80	18.73	31.41	7.60	15.00	19.12	
Ethylene diamine D *0.5 A *1.2 H ₂ O * 2.0HCl	32.12	7.92	18.73	18.96	31.90	9.19	18.08	19.11	
D,L-Aspartic Acid D *1.0 A *1.8 H ₂ O * 0.4HOAc	37.60	7.20	13.70		37.59	7.66	13.73		
D-Glutamic Acid D *1.0 A *1.8 H ₂ O * 0.3HOAc	39.18	7.45	13.44		39.47	7.52	13.29		
Squaric Acid D *1.0A *0.5 H ₂ O	42.10	5.89	12.27		9.37	42.29	5.72	12.73	9.42
D *0.5 A *0.75 H ₂ O	41.44	6.78	14.50		11.06	41.43	6.56	14.16	10.85

Fumaric Acid D *1.0 A *2.5 H ₂ O *2.5 EtOH	41.20	8.34	8.48		41.99	8.33	8.58	
1-hydroxy-2-naphthoic Acid D *1.2 A *1.0 H ₂ O * 1.0 EtOH	54.72	6.85	8.25		54.32	6.08	8.31	
1-hydroxy-2- naphthenesulfonic Acid D *0.65 A *2.4 H ₂ O	43.42	6.58	10.48		51.01	6.53	10.39	
2-Carboxyethyl phosphonic Acid D *1.1A *1.25 H ₂ O *1.0 CaCl ₂	29.08	5.87	9.00		29.34	5.83	8.70	
D *1.5 A *1.0 H ₂ O D *1.5 A *0.75 H ₂ O * 1.0 LiCl	33.15	6.49	9.64		33.17	6.56	9.28	
	29.65	5.77	8.30		29.66	5.71	8.84	

Phosphonoacetic Acid							
D *0.5 A *2.5 H ₂ O	32.58	6.83	12.66		32.16	6.73	13.33
D *2.0 A *1.0 H ₂ O	27.86	6.12	8.12		27.76	5.56	8.40
Phenyl phosphonic Acid							
D *1.0 A *0.5 H ₂ O	43.52	6.52	10.87		43.90	6.78	10.12
L-pyroglutamic Acid							
D *1.0 A *1.2 H ₂ O	42.20	7.19	15.15		42.35	7.10	15.01
HPF ₆							
D *1.0 A *0.5 H ₂ O	26.31	4.97	11.23		26.63	5.10	10.64

Biological Data

5 Some or all of the following assays are used to demonstrate the nitric oxide synthase inhibitory activity of the invention's compounds as well as demonstrate the useful pharmacological properties.

Citrulline Assay for Nitric Oxide Synthase

10

Nitric oxide synthase (NOS) activity can be measured by monitoring the conversion of L-[2,3-³H]-arginine to L-[2,3-³H]-citrulline (Bredt and Snyder, Proc. Natl. Acad. Sci. U.S.A., 87, 682-685, 1990 and Moore et al, J. Med. Chem., 39, 669-672, 1996). Human inducible NOS (hiNOS), human endothelial constitutive NOS (hecNOS) and human neuronal constitutive NOS (hncNOS) are each cloned from RNA extracted from human tissue. The cDNA for human inducible NOS (hiNOS) is isolated from a λcDNA library made from RNA extracted from a colon sample from a patient with ulcerative colitis. The cDNA for human endothelial constitutive NOS (hecNOS) is isolated from a λcDNA library made from RNA extracted from human umbilical vein endothelial cells (HUVEC) and the cDNA for human neuronal constitutive NOS (hncNOS) is isolated from a λcDNA library made from RNA extracted from human cerebellum obtained from a cadaver. The recombinant enzymes are expressed in Sf9 insect cells using a baculovirus vector (Rodi et al, in The Biology of Nitric Oxide, Pt. 4: Enzymology, Biochemistry and Immunology; Moncada, S., Feelisch, M., Busse, R., Higgs, E., Eds.; Portland Press Ltd.: London, 1995; pp 447-450). Enzyme activity is isolated from soluble cell extracts and partially purified by DEAE-Sepharose chromatography. To measure NOS activity, 10 μL of enzyme is added to 40 μL of 50 mM Tris (pH 7.6) in the presence or absence of test compounds and the reaction initiated by the addition of 50 μL of a reaction mixture containing 50mM Tris (pH 7.6), 2.0 mg/mL bovine serum albumin, 2.0 mM DTT, 4.0 mM CaCl₂, 20 μM FAD, 100 μM tetrahydrobiopterin, 0.4 mM NADPH and 60 μM L-arginine containing 0.9 μCi of L-[2,3-³H]-arginine. The final concentration of L-arginine in the assay is 30 μM. For hecNOS or

hncNOS, calmodulin is included at a final concentration of 40-100 nM. Following incubation at 37°C for 15 minutes, the reaction is terminated by addition of 400 µL of a suspension (1 part resin, 3 parts buffer) of Dowex 50W X-8 cation exchange resin in a stop buffer containing 10 mM EGTA, 100 mM HEPES, pH 5.5 and 1 mM L-citrulline. After 5 mixing the resin is allowed to settle and L-[2,3-³H]-Citrulline formation is determined by counting aliquots of the supernatant with a liquid scintillation counter. Results are reported in Table 1 as the IC₅₀ values of compounds for hiNOS, hecNOS and hncNOS.

10 In Vivo Assay

Rats can be treated with an intraperitoneal injection of 1-12.5 mg/kg of endotoxin (LPS) with or without oral administration of the nitric oxide synthase inhibitors. Plasma nitrite/nitrate levels can be determined 5 hours post-treatment. The results can be used to show that the administration of the nitric oxide synthase inhibitors decreases the rise in 15 plasma nitrite/nitrate levels, a reliable indicator of the production of nitric oxide induced by endotoxin.

Raw Cell Nitrite Assay

RAW 264.7 cells can be plated to confluence on a 96-well tissue culture plate grown 20 overnight (17h) in the presence of LPS to induce NOS. A row of 3-6 wells can be left untreated and served as controls for subtraction of nonspecific background. The media can be removed from each well and the cells washed twice with Kreb-Ringers-Hepes (25 mM, pH 7.4) with 2 mg/ml glucose. The cells are then placed on ice and incubated with 50 µL of buffer containing L-arginine (30 µM) +/- inhibitors for 1h. The assay can be initiated by 25 warming the plate to 37° C in a water bath for 1h. Production of nitrite by intracellular iNOS will be linear with time. To terminate the cellular assay, the plate of cells can be placed on ice and the nitrite-containing buffer removed and analyzed for nitrite using a previously published fluorescent determination for nitrite. T. P. Misko et al, Analytical Biochemistry, 214, 11-16 (1993).

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Human cartilage explant assay

Bone pieces are rinsed twice with Dulbecco's Phosphate Buffered Saline (GibcoBRL) and once with Dulbecco's Modified Eagles Medium (GibcoBRL) and placed into a petri dish with phenol red free Minimum Essential Medium (MEM) (GibcoBRL). Cartilage was cut into small explants of approximately 15-45 mg in weight and one or two

5 explants per well are placed into either 96 or 48 well culture plates with 200-500 μ L of culture media per well. The culture media was either a custom modification of Minimum Essential Medium(Eagle) with Earle's salts (GibcoBRL) prepared without L-Arginine, without L-Glutamine and without phenol red or a custom modification of serumless Neuman and Tytell (GibcoBRL) medium prepared without L-arginine, without insulin,

10 without ascorbic acid, without L-glutamine and without phenol red. Both are supplemented before use with 100 μ M L-Arginine (Sigma), 2 mM L-glutamine, 1X HL-1 supplement (BioWhittaker), 50 mg/ml ascorbic acid (Sigma) and 150 pg/ml recombinant human IL-1 β (RD Systems) to induce nitric oxide synthase. Compounds are then added in 10 μ L aliquots and the explants incubated at 37° C with 5% CO₂ for 18-24 hours. The day old supernatant

15 is then discarded and replaced with fresh culture media containing recombinant human IL-1 β and compound and incubated for another 20-24 hours. This supernatant is analyzed for nitrite with a fluorometric assay (Misko et al, *Anal. Biochem.*, 214, 11-16, 1993). All samples are done in quadruplicate. Unstimulated controls are cultured in media in the absence of recombinant human IL-1 β . IC₅₀ values (Table 1) are determined from plotting the

20 percent inhibition of nitrite production at six different concentrations of inhibitor.

Table 9 shows examples of biological activity for some of the compounds of the present invention.

25 Table 9. Biological Activity. Values represent averages across all experiments and all lots studied.

Example Number of Compound	hiNOS IC ₅₀ (μ M)	hecNOS IC ₅₀ (μ M)	hncNOS IC ₅₀ (μ M)	Human Cartilage IC ₅₀ (μ M)
Example 1	3.1	77	15	0.7
Example 2	4.4	302	58	8.2
Example 3	74	266	86	
Example 4	197	1100	539	
Example 7	3.4	78	17	
Example 11	0.9	26	6.0	

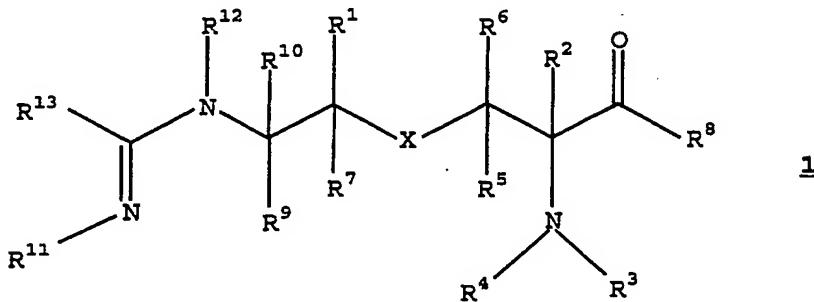
Example 16	7.2	>100	36	0.7
Example 18	12	>100	181	
Example 19	12	1080	220	

From the foregoing description, one skilled in the art can easily ascertain the essential characteristics of this invention, and without departing from the spirit and scope thereof, can
5 make various changes and modifications of the invention to adapt it to various usages and conditions.

What is claimed is:

1. A compound or a pharmaceutically acceptable salt thereof, the compound having a structure corresponding to Formula 1:

5



wherein:

X is selected from the group consisting of -S-, -S(O)-, and -S(O)2-;

10 R² is selected from the group consisting of C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₅ alkoxy-C₁ alkyl, and C₁-C₅ alkylthio-C₁ alkyl;
with respect to R³ and R⁸:

15 R⁸ is selected from the group consisting of -OR¹⁴ and -N(R¹⁵)(R¹⁶); and R³ is selected from the group consisting of -H, -OH, -C(O)-R¹⁷, -C(O)-O-R¹⁸, and -C(O)-S-R¹⁹; or

R⁸ is -N(R²⁰)-, and R³ is -C(O)-, wherein R⁸ and R³ together with the atoms to which they are attached form a ring; or

R⁸ is -O-, and R³ is -C(R²¹)(R²²)-, wherein R⁸ and R³ together with the atoms to which they are attached form a ring;

20 if R³ is -C(R²¹)(R²²)-, then R⁴ is -C(O)-O-R²³; otherwise R⁴ is -H;
R¹, R⁵, R⁶, and R⁷ independently are selected from the group consisting of -H, halogen, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, and C₁-C₅ alkoxy-C₁ alkyl;

R⁹ and R¹⁰ independently are selected from the group consisting of -H, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, and C₁-C₅ alkoxy-C₁ alkyl;

with respect to R¹¹ and R¹²:

R¹¹ is selected from the group consisting of -H, -OH, -C(O)-O-R²⁴, and -C(O)-S-R²⁵; and R¹² is selected from the group consisting of -H, -OH, -C(O)-O-R²⁶, and -C(O)-S-R²⁷; or

5 R¹¹ is -O-, and R¹² is -C(O)-, wherein R¹¹ and R¹² together with the atoms to which they are attached form a ring; or

R¹¹ is -C(O)-, and R¹² is -O-, wherein R¹¹ and R¹² together with the atoms to which they are attached form a ring; and

R¹³ is C₁ alkyl;

10 R¹⁴ is selected from the group consisting of -H and C₁-C₆ alkyl; wherein

when R¹⁴ is C₁-C₆ alkyl, R¹⁴ is optionally substituted by one or more moieties selected from the group consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl; with respect to R¹⁵ and R¹⁶:

15 R¹⁵ is selected from the group consisting of -H, alkyl, and alkoxy; and R¹⁶ is selected from the group consisting of -H, -OH, alkyl, alkoxy, -C(O)-R^{27a}, -C(O)-O-R²⁸, and -C(O)-S-R²⁹; wherein when R¹⁵ and R¹⁶ independently are alkyl or alkoxy, R¹⁵ and R¹⁶ independently are optionally substituted with one or more moieties selected from the group consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl; or

20 R¹⁵ is -H; and R¹⁶ is selected from the group consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl;

R¹⁷, R¹⁸, R¹⁹, R²⁰, R²¹, R²², R²³, R²⁴, R²⁵, R²⁶, R²⁷, R^{27a}, R²⁸, and R²⁹ independently are selected from the group consisting of -H and alkyl, wherein alkyl is optionally substituted by one or more moieties selected from the group consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl; and

25 when any of R¹, R², R³, R⁴, R⁵, R⁶, R⁷, R⁸, R⁹, R¹⁰, R¹¹, R¹², R¹³, R¹⁴, R¹⁵, R¹⁶, R¹⁷, R¹⁸, R¹⁹, R²⁰, R²¹, R²², R²³, R²⁴, R²⁵, R²⁶, R²⁷, R^{27a}, R²⁸, and R²⁹ independently is a moiety selected from the group consisting of alkyl, alkenyl, alkynyl, alkoxy, alkylthio, cycloalkyl, heterocyclyl, aryl, and heteroaryl, then the moiety is optionally

substituted by one or more substituent selected from the group consisting of -OH, alkoxy, and halogen.

2. The compound of Claim 1 wherein R⁸ is -OH.
5
3. The compound of Claim 2 wherein X is S.
4. The compound of Claim 3 wherein R³, R⁴, R¹¹ and R¹² each is -H.
- 10 5. The compound of Claim 4 wherein R¹, R⁵, R⁶, R⁷, R⁹, and R¹⁰ independently are selected from the group consisting of -H and C₁-C₃ alkyl.
6. The compound of Claim 5 wherein R⁵, R⁶, R⁷, R⁹, R¹⁰ each is -H.
- 15 7. The compound of Claim 6 wherein R¹³ is fluoromethyl.
8. The compound of Claim 6 wherein R¹³ is methyl.
9. The compound of Claim 8 wherein R² is C₁-C₆ alkyl optionally substituted with a
20 substituent selected from the group consisting of -OH, alkoxy, and halogen.
10. The compound of Claim 9 wherein R² is C₁ alkyl optionally substituted with halogen.
- 25 11. The compound of Claim 10 wherein R² is fluoromethyl.
12. The compound of Claim 10 wherein R² is methyl.
13. The compound of Claim 9 wherein R² is hydroxymethyl.

14. The compound of Claim 9 wherein R² is methoxymethyl.
15. The compound of Claim 8 wherein R¹ is selected from the group consisting of -H and C₁-C₆ alkyl optionally substituted with a substituent selected from the group consisting of -OH, alkoxy, and halogen.
5
16. The compound of Claim 15 wherein R¹ is -H.
17. The compound of Claim 15 wherein R¹ is C₁-C₆ alkyl optionally substituted with a substituent selected from the group consisting of alkoxy and halogen.
10
18. The compound of 17 wherein R¹ is ethyl.
19. The compound of Claim 15 wherein R¹ is C₁ alkyl optionally substituted with a substituent selected from the group consisting of -OH, alkoxy, and halogen.
15
20. The compound of Claim 19 wherein R¹ is methyl.

21. The compound of Claim 19 wherein R¹ is fluoromethyl.
20
22. The compound of Claim 19 wherein R¹ is hydroxymethyl.

23. The compound of Claim 19 wherein R¹ is C₁-C₆ alkoxyethyl.

- 25 24. The compound of Claim 19 wherein R¹ is C₁-C₆ methoxymethyl.

25. The compound of Claim 1 wherein R⁸ is -OR¹⁴.

26. The compound of Claim 25 wherein R¹⁴ is C₁-C₆ alkyl optionally substituted by one or more moieties selected from the group consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl.
- 5 27. The compound of Claim 26 wherein R¹⁴ is C₁-C₃ alkyl.
28. The compound of Claim 27 wherein R¹⁴ is methyl.
29. The compound of Claim 1 wherein R⁸ is -N(R¹⁵)(R¹⁶).
- 10 30. The compound of Claim 1 wherein R⁸ is -N(R²⁰)-, and R³ is -C(O)-, wherein R⁸ and R³ together with the atoms to which they are attached form a ring.
- 15 31. The compound of Claim 1 wherein R⁸ is -O-, and R³ is -C(R²¹)(R²²)-, wherein R⁸ and R³ together with the atoms to which they are attached form a ring.
32. The compound of Claim 1 wherein R¹¹ is selected from the group consisting of -OH, -C(O)-O-R²⁴, and -C(O)-S-R²⁵.
- 20 33. The compound of Claim 32 wherein R¹¹ is -OH.
34. The compound of Claim 33 wherein R¹² is -H.
35. The compound of Claim 1 wherein R¹¹ is -O-, and R¹² is -C(O)-, wherein R¹¹ and R¹² together with the atoms to which they are attached form a ring.
- 25 36. The compound of Claim 1 wherein R¹¹ is -C(O)-, and R¹² is -O-, wherein R¹¹ and R¹² together with the atoms to which they are attached form a ring.

37. The compound of Claim 1 wherein R¹² is selected from the group consisting of -OH, -C(O)-O-R²⁶, and -C(O)-S-R²⁷.

38. The compound of Claim 37 wherein R¹¹ is -H.

5 39. The compound of Claim 1 wherein the compound is in the form of a pharmaceutically-acceptable salt.

10 40. The pharmaceutically-acceptable salt of Claim 39 having at least one anionic counterion.

15 41. The pharmaceutically-acceptable salt of Claim 40 wherein the anionic counterion is selected from the group consisting of a halide, a carboxylate, a sulfonate, a sulfate, a phosphate, a phosphonate, a resin-bound anion, and a nitrate.

42. The pharmaceutically-acceptable salt of Claim 41 wherein the anionic counterion is a halide.

20 43. The pharmaceutically-acceptable salt of Claim 42 wherein the halide is chloride.

44. The pharmaceutically-acceptable salt of Claim 41 wherein the anionic counterion is a carboxylate.

25 45. The pharmaceutically-acceptable salt of Claim 44 wherein the carboxylate is selected from the group consisting of formate, acetate, propionate, trifluoroacetate, succinate, salicylate, DL-aspartate, D-aspartate, L-aspartate, DL-glutamate, D-glutamate, L-glutamate, glycerate, succinate, steric, DL-tartarate, D-tartarate, L-tartarate, (±)-mandelate, (R)-(-)-mandelate, (S)-(+) -mandelate, citrate, mucate, maleate, malonate, benzoate, DL-malate, D-malate, L-malate, hemimalate, 1-adamantaneacetate, 1-adamantanecarboxylate, flavianate, sulfonoacetate, (±)-lactate, L-(+)-lactate, D-(+)-lactate, pamoate, D-alpha-

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galacturonate, glycerate, DL-cystate, D-cystate, L-cystate, DL-homocystate, D-homocystate, L-homocystate, DL-cysteate, D-cysteate, L-cysteate, (4S)-hydroxy-L-proline, cyclopropane-1,1-dicarboxylate, 2,2-dimethylmalonate, squarate, tyrosine anion, proline anion, fumarate, 1-hydroxy-2-naphthoate, phosphonoacetate, carbonate, bicarbonate, 3-phosphonopropionate, DL-pyroglutamate, D-pyroglutamate, and L-pyroglutamate.

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46. The pharmaceutically-acceptable salt of Claim 41 wherein the anionic counterion is a sulfonate.

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47. The pharmaceutically-acceptable salt of Claim 46 wherein the sulfonate is selected from the group consisting of methanesulfonate, toluenesulfonate, benzenesulfonate, trifluoromethylsulfonate, ethanesulfonate, (\pm)-camphorsulfonate, naphthalenesulfonate, 1R-(-)-camphorsulfonate, 1S-(+)-camphorsulfonate, 2-mesitylenesulfonate, 1,5-naphthalenedisulfonate, 1,2-ethanedisulfonate, 1,3-propanedisulfonate, 3-(N-morpholino)propane sulfonate, biphenylsulfonate, isethionate, and 1-hydroxy-2-naphthalenesulfonate.

15

48. The pharmaceutically-acceptable salt of Claim 41 wherein the anionic counterion is a sulfate.

20

49. The pharmaceutically-acceptable salt of Claim 48 wherein the sulfate is selected from the group consisting of sulfate, monopotassium sulfate, monosodium sulfate, and hydrogen sulfate.

25

50. The pharmaceutically-acceptable salt of Claim 41 wherein the anionic counterion is a sulfamate.

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51. The pharmaceutically-acceptable salt of Claim 41 wherein the anionic counterion is a phosphate.

52. The pharmaceutically-acceptable salt of Claim 51 wherein the phosphate is selected from the group consisting of phosphate, dihydrogen phosphate, potassium hydrogen phosphate, dipotassium phosphate, potassium phosphate, sodium hydrogen phosphate, disodium phosphate, sodium phosphate, calcium phosphate, and hexafluorophosphate.

53. The pharmaceutically-acceptable salt of Claim 41 wherein the anionic counterion is a phosphonate.

10 54. The pharmaceutically-acceptable salt of Claim 53 wherein the phosphonate is selected from the group consisting of vinylphosphonate, 2-carboxyethylphosphonate and phenylphosphonate.

15 55. The pharmaceutically-acceptable salt of Claim 41 wherein the anionic counterion is a resin-bound anion.

56. The pharmaceutically-acceptable salt of Claim 55 wherein the resin-bound anion is selected from the group consisting of a resin comprising polyacrylate and a resin comprising sulfonated poly(styrene divinylbenzene).

20 57. The pharmaceutically-acceptable salt of Claim 41 wherein the anionic counterion is nitrate.

58. The pharmaceutically-acceptable salt of Claim 40 wherein the anion is selected from the group consisting of DL-ascorbate, D-ascorbate, and L-ascorbate.

25 59. The pharmaceutically-acceptable salt of Claim 39 having at least one cationic counterion.

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60. The pharmaceutically-acceptable salt of Claim 59 wherein the cationic counterion is selected from the group consisting of an ammonium cation, a alkali metal cation, an alkaline earth metal cation, a transition metal cation, and a resin-bound cation.

5

61. The pharmaceutically-acceptable salt of Claim 60 wherein the cationic counterion is an ammonium cation.

10

62. The pharmaceutically-acceptable salt of Claim 61 wherein the ammonium cation is selected from the group consisting of ammonium, methyl ammonium, dimethylammonium, trimethylammonium, tetramethylammonium, ethanolammonium, dicyclohexylammonium, guanidinium, and ethylenediammonium cation.

15 63. The pharmaceutically-acceptable salt of Claim 60 wherein the cationic counterion is an alkali metal cation.

20

64. The pharmaceutically-acceptable salt of Claim 63 wherein the alkali metal cation is selected from the group consisting of lithium cation, sodium cation, potassium cation, and cesium cation.

65. The pharmaceutically-acceptable salt of Claim 60 wherein the cationic counterion is an alkaline earth metal cation.

25 66. The pharmaceutically-acceptable salt of Claim 65 wherein the alkaline earth metal cation is selected from the group consisting of beryllium cation, magnesium cation, and calcium cation.

30

67. The pharmaceutically-acceptable salt of Claim 60 wherein the cationic counterion is a transition metal cation.

68. The pharmaceutically-acceptable salt of Claim 67 wherein the transition metal cation is a zinc cation.

69. The pharmaceutically-acceptable salt of Claim 60 wherein the cationic counterion is a resin-bound cation.

5 70. The pharmaceutically-acceptable salt of Claim 69 wherein the resin-bound cation is a cationically functionalized poly(styrene divinylbenzene) resin.

10 71. The pharmaceutically-acceptable salt of Claim 70 wherein the resin-bound cation is an aminated poly(styrene divinylbenzene) resin.

72. The pharmaceutically-acceptable salt of Claim 69 wherein the resin-bound cation is a cationically functionalized polyacrylic resin.

15 73. The pharmaceutically-acceptable salt of Claim 69 wherein the resin-bound cation is an aminated polyacrylic resin.

74. *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-methyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

20 75. *S*-[(1*S*)-2-[(1-Iminoethyl)amino]-1-methylethyl]-2-methyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

25 76. *S*-[(1*R/S*)-2-[(1-Iminoethyl)amino]-1-ethylethyl]-2-methyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

77. *S*-[(1*S*)-2-[(1-iminoethyl)amino]-1-(fluoromethyl)ethyl]-2-methyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

30

78. *S*-[2-[(1-Iminoethyl)amino]ethyl]-2-ethyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

79. *S*-[(1*R*)-2-[(1-Iminoethyl)amino]-1-methylethyl]-2-methyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

80. *S*-[(1*R*)-2-[(1-Iminoethyl)amino]-1-methylethyl]-2-ethyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

10 81. *S*-[(1*R/S*)-2-[(1-Iminoethyl)amino]-1-ethylethyl]-2-ethyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

82. *S*-[(1*S*)-2-[(1-Iminoethyl)amino]-1-fluoromethylethyl]-2-ethyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

15 82. 2-[[[2-(1-Iminoethyl)amino]ethyl]thio]methyl]-D-valine, or a pharmaceutically acceptable salt thereof.

20 84. 2-[[[(1*R*)-2-[(1-Iminoethyl)amino]-1-methylethyl]thio]-methyl]-D-valine, or a pharmaceutically acceptable salt thereof.

85. 2-[[[(1*R/S*)-2-(1-Iminoethyl)amino)-1-ethylethyl]-thio]methyl]-D-valine, or a pharmaceutically acceptable salt thereof.

25 86. 2-[[[(1*S*)-2-[(1-Iminoethyl)amino]-1-fluoromethylethyl]-thio]methyl]-D-valine, or a pharmaceutically acceptable salt thereof.

87. *S*-[(*R/S*)-2-[(1-Iminoethyl)amino]-1-(trifluoromethyl)ethyl]-2-methyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

88. S-[2-(1-Iminoethylamino)ethyl]-2-methyl-(D/L)-cysteine, or a pharmaceutically acceptable salt thereof.

89. S-[2-[(1-Iminoethyl)amino]ethyl]-2-fluoromethyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

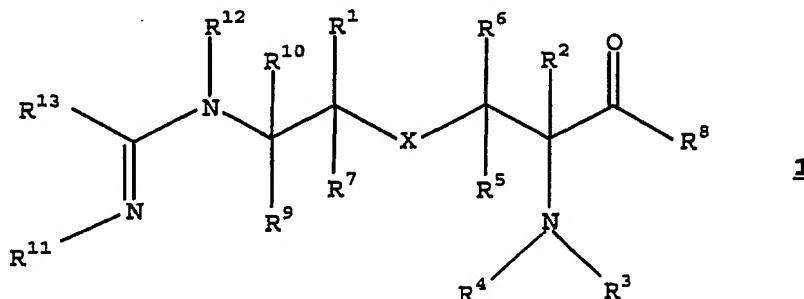
90. (2R)-2-Amino-3-[[2-[(1-iminoethyl)amino]ethyl]sulfinyl]-2-methylpropanoic acid, or a pharmaceutically acceptable salt thereof.

10 91. (2R)-2-Amino-3-[[2-[(1-iminoethyl)amino]ethyl]sulfonyl]-2-methylpropanoic acid, or a pharmaceutically acceptable salt thereof.

92. N-{4-Chlorophenyl)methylene]-S-[2-[(4-chlorophenyl)methylene]amino]ethyl]-L-cysteine, methyl ester.

15 93. N-[4-Chlorophenyl)methylene]-S-[2-[(4-chlorophenyl)methylene]amino]ethyl]-2-methyl-D/L-cysteine, methyl ester.

94. A method for the treatment or prevention of an inflammation-related disorder
20 wherein the method comprises treating a subject in need thereof with an inflammation-related disorder-treating or preventing amount of a compound or a pharmaceutically acceptable salt thereof, the compound having a structure corresponding to Formula 1:



wherein:

X is selected from the group consisting of -S-, -S(O)-, and -S(O)₂-;

R² is selected from the group consisting of C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₅ alkoxy-C₁ alkyl, and C₁-C₅ alkylthio-C₁ alkyl;

5 with respect to R³ and R⁸:

R⁸ is selected from the group consisting of -OR¹⁴ and -N(R¹⁵)(R¹⁶); and R³ is selected from the group consisting of -H, -OH, -C(O)-R¹⁷, -C(O)-O-R¹⁸, and -C(O)-S-R¹⁹; or

10 R⁸ is -N(R²⁰)-, and R³ is -C(O)-, wherein R⁸ and R³ together with the atoms to which they are attached form a ring; or

R⁸ is -O-, and R³ is -C(R²¹)(R²²)-, wherein R⁸ and R³ together with the atoms to which they are attached form a ring;

if R³ is -C(R²¹)(R²²)-, then R⁴ is -C(O)-O-R²²; otherwise R⁴ is -H;

15 R¹, R⁵, R⁶, and R⁷ independently are selected from the group consisting of -H, halogen, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, and C₁-C₅ alkoxy-C₁ alkyl;

R⁹ and R¹⁰ independently are selected from the group consisting of -H, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, and C₁-C₅ alkoxy-C₁ alkyl;

with respect to R¹¹ and R¹²:

20 R¹¹ is selected from the group consisting of -H, -OH, -C(O)-O-R²⁴, and -C(O)-S-R²⁵; and R¹² is selected from the group consisting of -H, -OH, -C(O)-O-R²⁶, and -C(O)-S-R²⁷; or

R¹¹ is -O-, and R¹² is -C(O)-, wherein R¹¹ and R¹² together with the atoms to which they are attached form a ring; or

25 R¹¹ is -C(O)-, and R¹² is -O-, wherein R¹¹ and R¹² together with the atoms to which they are attached form a ring; and

R¹³ is C₁ alkyl;

R^{14} is selected from the group consisting of -H and C_1 - C_6 alkyl; wherein when R^{14} is C_1 - C_6 alkyl, R^{14} is optionally substituted by one or more moieties selected from the group consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl; with respect to R^{15} and R^{16} :

5 R^{15} is selected from the group consisting of -H, alkyl, and alkoxy; and R^{16} is selected from the group consisting of -H, -OH, alkyl, alkoxy, -C(O)- R^{27a} , -C(O)-O- R^{28} , and -C(O)-S- R^{29} ; wherein when R^{15} and R^{16} independently are alkyl or alkoxy, R^{15} and R^{16} independently are optionally substituted with one or more moieties selected from the group consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl; or

10 R^{15} is -H; and R^{16} is selected from the group consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl;

15 R^{17} , R^{18} , R^{19} , R^{20} , R^{21} , R^{22} , R^{23} , R^{24} , R^{25} , R^{26} , R^{27} , R^{27a} , R^{28} , and R^{29} independently are selected from the group consisting of -H and alkyl, wherein alkyl is optionally substituted by one or more moieties selected from the group consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl; and

20 when any of R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^9 , R^{10} , R^{11} , R^{12} , R^{13} , R^{14} , R^{15} , R^{16} , R^{17} , R^{18} , R^{19} , R^{20} , R^{21} , R^{22} , R^{23} , R^{24} , R^{25} , R^{26} , R^{27} , R^{27a} , R^{28} , and R^{29} independently is a moiety selected from the group consisting of alkyl, alkenyl, alkynyl, alkoxy, alkylthio, cycloalkyl, heterocyclyl, aryl, and heteroaryl, then the moiety is optionally substituted by one or more substituent selected from the group consisting of -OH, alkoxy, and halogen.

95. The method of Claim 94 wherein the compound is *S*-[2-[(1-iminoethyl)amino]ethyl]-2-methyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

96. The method of Claim 94 wherein the compound is 2-[[2-[(1-iminoethyl)amino]ethyl]thio]methyl]-*O*-methyl-D-serine, or a pharmaceutically acceptable salt thereof.

97. The method of Claim 94 wherein the compound is *S*-[(1*S*)-2-[(1-iminoethyl)amino]-1-methylethyl]-2-methyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

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98. The method of Claim 94 wherein the compound is *S*-[(1*R/S*)-2-[(1-iminoethyl)amino]-1-ethylethyl]-2-methyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

10 99. The method of Claim 94 wherein the compound is *S*-[(1*S*)-2-[(1-iminoethyl)amino]-1-(fluoromethyl)ethyl]-2-methyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

15 100. The method of Claim 94 wherein the compound is *S*-[2-[(1-iminoethyl)amino]ethyl]-2-ethyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

101. The method of Claim 94 wherein the compound is *S*-[(1*R*)-2-[(1-iminoethyl)amino]-1-methylethyl]-2-methyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

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102. The method of Claim 94 wherein the compound is *S*-[(1*R*)-2-[(1-iminoethyl)amino]-1-methylethyl]-2-ethyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

25

103. The method of Claim 94 wherein the compound is *S*-[(1*R/S*)-2-[(1-iminoethyl)amino]-1-ethylethyl]-2-ethyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

104. The method of Claim 94 wherein the compound is *S*-[(1*S*)-2-[(1-iminoethyl)amino]-1-fluoromethylethyl]-2-ethyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

5 105. The method of Claim 94 wherein the compound is 2-[[[2-(1-iminoethyl)amino]ethyl]thio]methyl]-D-valine, or a pharmaceutically acceptable salt thereof.

10 106. The method of Claim 94 wherein the compound is 2-[[[(1*R*)-2-[(1-iminoethyl)amino]-1-methylethyl]thio]-methyl]-D-valine, or a pharmaceutically acceptable salt thereof.

15 107. The method of Claim 94 wherein the compound is 2-[[[(1*R/S*)-2-(1-iminoethyl)amino]-1-ethylethyl]-thio]methyl]-D-valine, or a pharmaceutically acceptable salt thereof.

108. The method of Claim 94 wherein the compound is 2-[[[(1*S*)-2-[(1-iminoethyl)amino]-1-fluoromethylethyl]-thio]methyl]-D-valine, or a pharmaceutically acceptable salt thereof.

20 109. The method of Claim 94 wherein the compound is *S*-[(*R/S*)-2-[(1-iminoethyl)amino]-1-(trifluoromethyl)ethyl]-2-methyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

25 110. The method of Claim 94 wherein the compound is *S*-[2-(1-iminoethylamino)ethyl]-2-methyl-(D/L)-cysteine, or a pharmaceutically acceptable salt thereof.

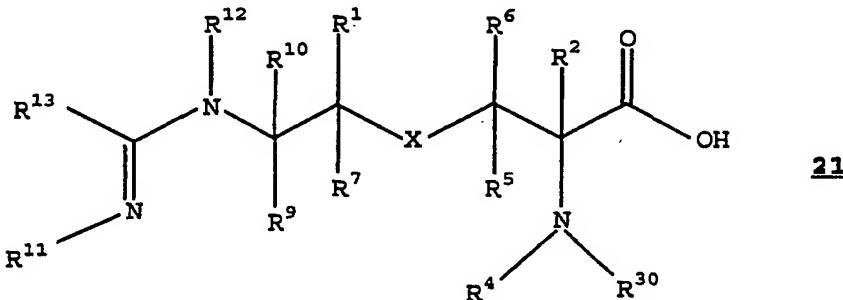
30 111. The method of Claim 94 wherein the compound is *S*-[2-[(1-iminoethyl)amino]ethyl]-2-fluoromethyl-L-cysteine, or a pharmaceutically acceptable salt thereof.

112. The method of Claim 94 wherein the compound is (2*R*)-2-amino-3-[[2-[(1-iminoethyl)amino]ethyl]sulfinyl]-2-methylpropanoic acid, or a pharmaceutically acceptable salt thereof.

5

113. The method of Claim 94 wherein the compound is (2*R*)-2-amino-3-[[2-[(1-iminoethyl)amino]ethyl]sulfonyl]-2-methylpropanoic acid, or a pharmaceutically acceptable salt thereof.

10 114. A method for the preparation of a compound or a pharmaceutically acceptable salt thereof, the compound having a structure corresponding to Formula 21:



15 or a salt thereof,

wherein:

X is selected from the group consisting of -S-, -S(O)-, and -S(O)₂-;

R² is selected from the group consisting of C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₅ alkoxy-C₁ alkyl, and C₁-C₅ alkylthio-C₁ alkyl;

20 R³⁰ is selected from the group consisting of -H, -OH, -C(O)-R¹⁷, -C(O)-O-R¹⁸, and -C(O)-S-R¹⁹;

R¹, R⁵, R⁶, and R⁷ independently are selected from the group consisting of -H, halogen, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, and C₁-C₅ alkoxy-C₁ alkyl;

R^9 and R^{10} independently are selected from the group consisting of -H, C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, and C_1 - C_5 alkoxy- C_1 alkyl;
with respect to R^{11} and R^{12} :

5 R^{11} is selected from the group consisting of -H, -OH, -C(O)-O-R²⁴, and -C(O)-S-R²⁵; and R^{12} is selected from the group consisting of -H, -OH, -C(O)-O-R²⁶, and -C(O)-S-R²⁷; or

10 R^{11} is -O-, and R^{12} is -C(O)-, wherein R^{11} and R^{12} together with the atoms to which they are attached form a ring; or

15 R^{11} is -C(O)-, and R^{12} is -O-, wherein R^{11} and R^{12} together with the atoms to which they are attached form a ring; and

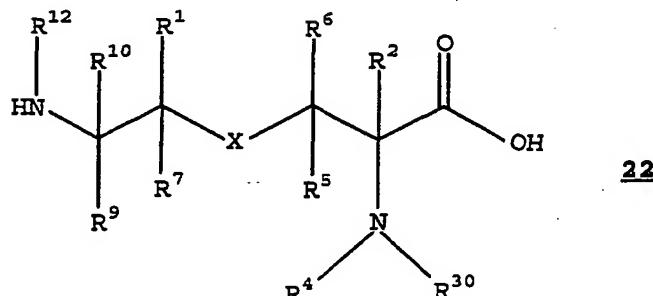
R^{13} is C_1 alkyl;

20 R^{17} , R^{18} , R^{19} , R^{24} , R^{25} , R^{26} , R^{27} , and R^{27a} independently are selected from the group consisting of -H and alkyl, which is optionally substituted by one or more moieties selected from the group consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl; and

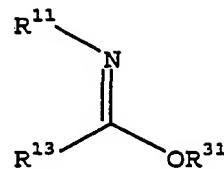
25 when any of R^1 , R^2 , R^4 , R^5 , R^6 , R^7 , R^9 , R^{10} , R^{11} , R^{12} , R^{13} , R^{14} , R^{15} , R^{16} , R^{17} , R^{18} , R^{19} , R^{24} , R^{25} , R^{26} , R^{27} , and R^{27a} independently is a moiety selected from the group consisting of alkyl, alkenyl, alkynyl, alkoxy, alkylthio, cycloalkyl, heterocyclyl, aryl, and heteroaryl, then the moiety is optionally substituted by one or more substituent

20 selected from the group consisting of -OH, alkoxy, and halogen;

wherein the method comprises treating a diamine compound having a structure corresponding to Formula 22:



or a pharmaceutically acceptable salt thereof, with an alkyl acetimidate having a structure corresponding to Formula 23:



23

5

or a salt thereof, wherein R³¹ is C₁-C₆ alkyl,

to produce the compound corresponding to Formula 21.

115. The method of Claim 114 wherein R¹¹ is selected from the group consisting of -H and -OH.

116. The method of claim 115 wherein R¹¹ is -H.

117. The method of Claim 115 wherein R¹¹ is -OH.

118. The method of Claim 115 wherein R¹³ is methyl or halomethyl.

119. The method of Claim 118 wherein R¹³ is methyl.

120. The method of Claim 115 wherein R³¹ is C₁-C₃ alkyl.

121. The method of Claim 120 wherein R³¹ is ethyl.

122. The method of Claim 114 wherein the treating of the diamine compound with the alkyl acetimidate compound is performed in the presence of a base.

123. The method of Claim 122 wherein the base is selected from the group consisting of a hydrazine, a metal sulfide, a metal hydroxide, a metal alkoxide, an amine, a hydroxylamine, a metal amide complex, and a basic resin.

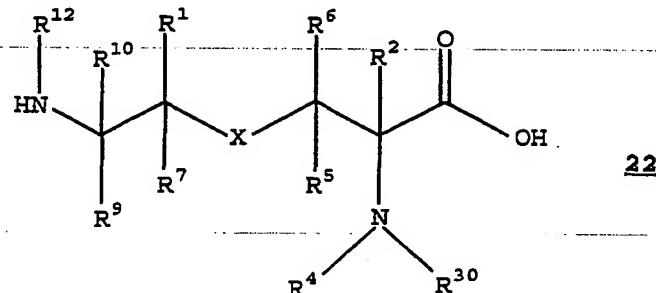
5 124. The method of Claim 123 wherein the base is a basic resin.

125. The method of Claim 124 wherein the basic resin is a polymer-bound diazabicyclo[4.4.0]dec-2-ene.

10 126. The method of Claim 123 wherein the base is an amine.

127. The method of Claim 126 wherein the base is selected from the group consisting of 1,5-diazabicyclo[4.3.0]non-5-ene; 1,4-diazabicyclo[2.2.2]octane; and 1,8-diazabicyclo[5.4.0]undec-7-ene.

15 128. A method for the preparation of a diamine compound having a structure corresponding to Formula 22:



20

or a pharmaceutically acceptable salt thereof, wherein:

X is selected from the group consisting of -S-, -S(O)-, and -S(O)₂;

R² is selected from the group consisting of C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₅ alkoxy-C₁ alkyl, and C₁-C₅ alkylthio-C₁ alkyl;

R^{30} is selected from the group consisting of -H, -OH, -C(O)-R¹⁷, -C(O)-O-R¹⁸, and -C(O)-S-R¹⁹;

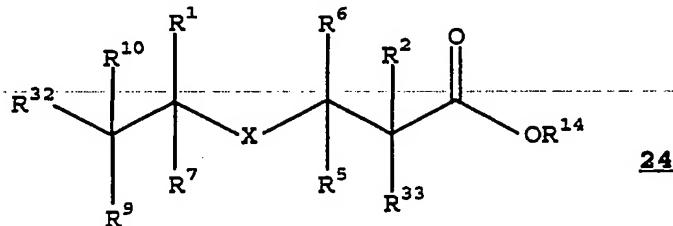
R^1 , R^5 , R^6 , and R^7 independently are selected from the group consisting of -H, halogen, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, and C₁-C₅ alkoxy-C₁ alkyl;

5 R^9 and R^{10} independently are selected from the group consisting of -H, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, and C₁-C₅ alkoxy-C₁ alkyl;

R^{12} is selected from the group consisting of -H, -OH, -C(O)-O-R²⁶, and -C(O)-S-R²⁷; and

10 when any of R^1 , R^2 , R^4 , R^5 , R^6 , R^7 , R^9 , R^{10} , R^{12} , R^{14} , R^{15} , R^{16} , R^{17} , R^{18} , R^{19} , R^{24} , R^{25} , R^{26} , R^{27} , and R^{27a} independently is a moiety selected from the group consisting of alkyl, alkenyl, alkynyl, alkoxy, alkylthio, cycloalkyl, heterocyclyl, aryl, and heteroaryl, then the moiety is optionally substituted by one or more substituent selected from the group consisting of -OH, alkoxy, and halogen;

15 wherein the method comprises treating a protected diamine compound having the structure corresponding to Formula 24:



20 or a pharmaceutically acceptable salt thereof, wherein R^{33} is selected from the group consisting of -NH₂ and a protected amino group; and R^{32} is a protected amino group; and R^{14} is selected from the group consisting of -H and C₁-C₆ alkyl; wherein when R^{14} is C₁-C₆ alkyl, R^{14} is optionally substituted by one or more moieties selected from the group consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl;

wherein the treating is performed with a deprotecting reagent, thereby producing the diamine compound.

129. The method of Claim 128 wherein R³³ is a 4-chlorobenzylimino group.
5
130. The method of Claim 129 wherein R³² is -NH₂.
131. The method of Claim 129 wherein R³² is a 4-chlorobenzylimino group.
- 10 132. The method of Claim 131 wherein R³³ is a 4-chlorobenzylimino group.
133. The method of Claim 134 wherein the treating of compound 24 with the deprotecting reagent is performed in the presence of water.
- 15 134. The method of Claim 133 wherein the deprotecting reagent is an acid.
135. The method of Claim 134 wherein the acid is selected from the group consisting of hydrochloric acid, hydrobromic acid, hydroiodic acid, sulfuric acid, methanesulfonic acid, trifluoromethanesulfonic acid, toluenesulfonic acid, 20 benzenesulfonic acid, trifluoroacetic acid, phosphoric acid, phosphorus acid, and acetic acid.
136. The method of Claim 128 wherein R³³ is a t-butoxycarbonylamino group.
25
137. The method of Claim 136 wherein R³² is -NH₂.
138. The method of Claim 136 wherein the deprotecting reagent is an acid.
139. The method of Claim 138 wherein the acid is selected from the group consisting 30 of hydrochloric acid, hydrobromic acid, hydroiodic acid, sulfuric acid, methanesulfonic acid, trifluoromethanesulfonic acid, toluenesulfonic acid, benzenesulfonic acid, trifluoroacetic acid, phosphoric acid, phosphorus acid, and acetic acid.

140. The method of Claim 138 wherein the treating of compound 24 with the deprotecting reagent is performed in the presence of water.

141. The method of Claim 128 wherein R³³ is an N-phthalimido group.

5 142. The method of Claim 141 wherein R³² is -NH₂.

143. The method of Claim 141 wherein the treating of compound 24 with the deprotecting reagent is performed in the presence of water.

10 144. The method of Claim 143 wherein the deprotecting reagent is selected from the group consisting of an acid and a base.

145. The method of Claim 144 wherein the deprotecting reagent is a base.

15 146. The method of Claim 145 wherein the base is selected from the group consisting of a hydrazine, a metal sulfide, a metal hydroxide, a metal alkoxide, an amine, a hydroxylamine, and a metal amide complex.

20 147. The method of Claim 146 wherein the deprotecting reagent is an acid.

148. The method of Claim 147 wherein the acid is selected from the group consisting of hydrochloric acid, hydrobromic acid, hydroiodic acid, sulfuric acid, methanesulfonic acid, trifluoromethanesulfonic acid, toluenesulfonic acid,

25 benzenesulfonic acid, trifluoroacetic acid, phosphoric acid, phosphorus acid, and acetic acid.

149. The method of Claim 148 wherein R¹⁴ is -H.

30 150. The method of Claim 149 wherein R³³ is -NH₂, or a salt thereof.

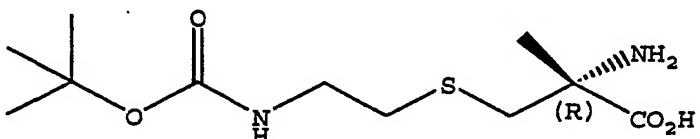
151. The method of Claim 150 wherein R^2 is methyl.

152. The method of Claim 151 wherein R^1 , R^5 , R^6 , R^7 , R^9 , and R^{10} each is -H.

5 153. The method of Claim 152 wherein R^{33} is a t-butoxycarbonylamino group.

154. The method of Claim 153 wherein the compound of Formula 24 has a structure corresponding to Formula 25:

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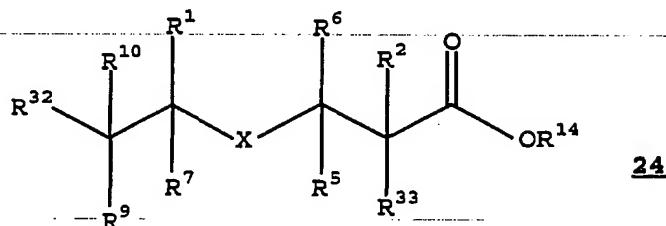


25

or a pharmaceutically-acceptable salt thereof.

15

155. A method for the preparation of a protected diamine compound having the structure corresponding to Formula 24:



24

or a pharmaceutically-acceptable salt thereof, wherein:

20

X is -S-;

R^2 is selected from the group consisting of C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_1 - C_5 alkoxy- C_1 alkyl, and C_1 - C_5 alkylthio- C_1 alkyl;

R^1 , R^5 , R^6 , and R^7 independently are selected from the group consisting of -H, halogen, C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, and C_1 - C_5 alkoxy- C_1 alkyl;

R⁹ and R¹⁰ independently are selected from the group consisting of -H, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, and C₁-C₅ alkoxy-C₁ alkyl;

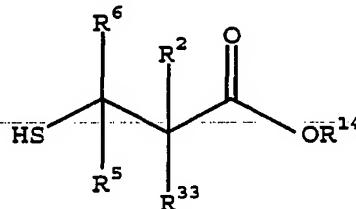
R³³ is selected from the group consisting of -NH₂ and a protected amino group; and R³² is a protected amino group;

5 R¹⁴ is selected from the group consisting of -H and C₁-C₆ alkyl; wherein when R¹⁴ is C₁-C₆ alkyl, R¹⁴ is optionally substituted by one or more moieties selected from the group consisting of cycloalkyl, heterocyclyl, aryl, and heteroaryl;

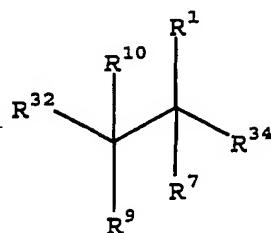
10 when any of R¹, R², R⁵, R⁶, R⁷, R⁹, R¹⁰, and R¹⁴ independently is a moiety selected from the group consisting of alkyl, alkenyl, alkynyl, alkoxy, alkylthio, cycloalkyl, heterocyclyl, aryl, and heteroaryl, then the moiety is optionally substituted by one or more substituent selected from the group consisting of -OH, alkoxy, and halogen;

wherein the method comprises treating a sulphydryl compound having the structure of Formula 26:

15

26

20

27

wherein R³⁴ is a nucleophilic substitution leaving group;
thereby forming the protected diamine compound.

156. The method of Claim 155 wherein the treating is performed in the presence of a
5 base.

157. The method of Claim 156 wherein the base is selected from the group consisting
of a hydrazine, a metal sulfide, a metal hydroxide, a metal alkoxide, an amine, a
hydroxylamine, a metal hydride, and a metal amide complex.

10 158. The method of Claim 157 wherein the base is an alkali metal hydroxide.

159. The method of Claim 158 wherein the base is selected from the group consisting
of potassium hydroxide and sodium hydroxide.

15 160. The method of Claim 157 wherein the base is a metal hydride.

161. The method of Claim 160 wherein the base is selected from the group consisting
of potassium hydride, sodium hydride, and calcium hydride.

20 162. The method of Claim 156 wherein R³⁴ is selected from the group consisting of
chloro, bromo, iodo, methanesulfonato, toluenesulfonato, benzenesulfonato, and
trifluoromethanesulfonato.

25 163. The method of Claim 162 wherein R³⁴ is chloro, bromo, and iodo.

164. The method of Claim 163 wherein R³⁴ is bromo.

165. The method of Claim 156 wherein R³³ is -NH₂.

30

166. The method of Claim 156 wherein R² is C₁-C₆ alkyl optionally substituted by one or more substituent selected from the group consisting of -OH, alkoxy and halogen.

5 167. The method of Claim 166 wherein R² is C₁-C₃ alkyl optionally substituted by one or more substituent selected from the group consisting of -OH, alkoxy and halogen.

10 168. The method of Claim 167 wherein R² is C₁-C₃ alkyl.

169. The method of Claim 168 wherein R² is methyl.

170. The method of Claim 156 wherein R⁵ and R⁶ each is -H.

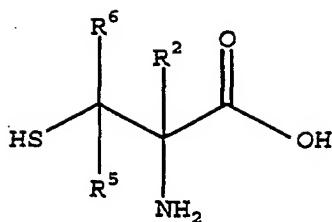
15 171. The method of Claim 156 wherein R¹ and R⁷ independently are selected from the group consisting of H and C₁-C₆ alkyl optionally substituted by one or more halogen.

172. The method of Claim 171 wherein R¹ and R⁷ each is -H.

20 173. The method of Claim 156 wherein R³² is selected from the group consisting of a 4-chlorobenzylimino group, a t-butoxycarbonylamino group, an N-phthalimido group, and a benzyloxycarbonylamino group.

25 174. The method of Claim 173 wherein R³² is a t-butoxycarbonylamino group.

175. A method for the preparation of a sulphydryl compound having the structure of Formula 28:

28

or a salt thereof, wherein:

R^2 is selected from the group consisting of C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_1 - C_5 alkoxy- C_1 alkyl, and C_1 - C_5 alkylthio- C_1 alkyl;

5

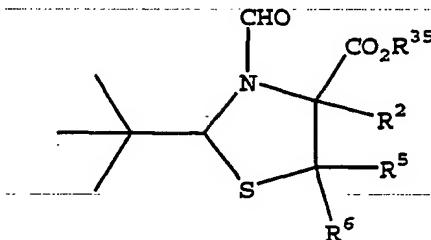
R^5 and R^6 independently are selected from the group consisting of -H, halogen, C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, and C_1 - C_5 alkoxy- C_1 alkyl;

10

when any of R^2 , R^5 , and R^6 independently is a moiety selected from the group consisting of alkyl, alkenyl, alkynyl, alkoxyalkyl, and alkylthioalkyl, then the moiety is optionally substituted by one or more substituent selected from the group consisting of -OH, alkoxy, and halogen;

wherein the method comprises treating under hydrolysis conditions a thiazolidine compound having the structure of Formula 29:

15

29

or a salt thereof, wherein R^{35} is a moiety selected from the group consisting of -H and C_1 - C_6 alkyl;

thereby forming the sulphydryl compound.

20

176. The method of Claim 175 wherein the hydrolysis conditions comprising contacting the thiazolidine compound with an acid in the presence of water.

177. The method of Claim 176 wherein the acid is selected from the group consisting of hydrochloric acid, hydrobromic acid, hydroiodic acid, sulfuric acid, methanesulfonic acid, trifluoromethanesulfonic acid, toluenesulfonic acid, benzenesulfonic acid, trifluoroacetic acid, phosphoric acid, phosphorus acid, and acetic acid.

10 178. The method of Claim 175 wherein R⁵ and R⁶ each is -H.

179. The method of Claim 175 wherein R² is C₁-C₆ alkyl optionally substituted by one or more substituent selected from the group consisting of -OH, alkoxy, and halogen.

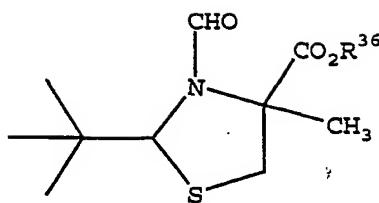
15 180. The method of Claim 179 wherein R² is C₁-C₃ alkyl optionally substituted by one or more substituent selected from the group consisting of alkoxy and halogen.

181. The method of Claim 180 wherein R² is C₁-C₃ alkyl.

20 182. The method of Claim 181 wherein R² is methyl.

183. The method of Claim 175 wherein R³⁵ is methyl.

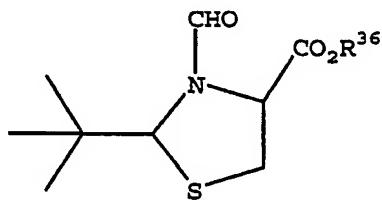
25 184. A method for the preparation of a methyl thiazolidine compound having the structure of Formula 30:



or a pharmaceutically-acceptable salt thereof;

wherein R³⁶ is C₁-C₆ alkyl;

5 wherein the method comprises treating under methylation conditions a deprotonatable thiazolidine compound having the structure of Formula 31:



10 thereby forming the methyl thiazolidine compound.

185. The method of Claim 184 wherein the methylation conditions comprise treating the deprotonatable thiazolidine compound with a base and a methylating agent.

15 186. The method of Claim 185 wherein the base is selected from the group consisting of a metal hydroxide, a metal alkoxide, a metal hydride, a metal alkyl, and a metal amide complex.

187. The method of Claim 186 wherein the base is a metal amide complex.

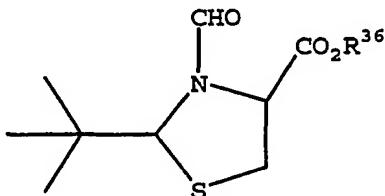
20 188. The method of Claim 187 wherein the base is selected from the group consisting of lithium hexamethyldisilazide, sodium hexamethyldisilazide, potassium hexamethyldisilazide, lithium diisopropyl amide, sodium diisopropyl amide, potassium diisopropyl amide, sodium amide, lithium amide, potassium amide,

sodium diethylamide, lithium diethylamide, potassium diethylamide, methyl lithium, t-butyl lithium, sec-butyl lithium, methyl sodium, t-butyl sodium, sec-butyl sodium, and methyl magnesium bromide.

5 189. The method of Claim 184 wherein R³⁶ is C₁-C₃ alkyl.

190. The method of Claim 189 wherein R³⁶ is methyl.

191. A method for the preparation of a deprotonatable thiazolidine compound having
10 the structure of Formula 31:



wherein R³⁶ is C₁-C₆ alkyl;

15 wherein the method comprises contacting under condensing conditions a cysteine C₁-C₆ alkyl ester with pivalaldehyde; thereby forming the deprotonatable thiazolidine compound.

192. The method of Claim 191 wherein the condensing conditions comprise
20 performing the contacting in the presence of a base.

193. The method of Claim 192 wherein the base is selected from the group consisting of a hydrazine, a metal sulfide, a metal hydroxide, a metal alkyl base, a metal alkoxide, an amine, a hydroxylamine, and a metal amide complex.

25 194. The method of Claim 193 wherein the base is a metal amide complex.

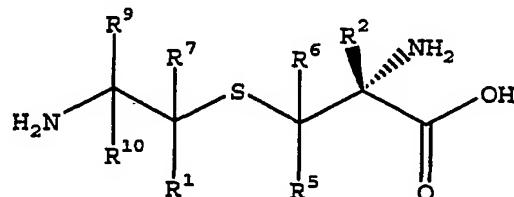
195. The method of Claim 194 wherein the base is lithium bis(trimethylsilyl)amide.

196. The method of Claim 191 wherein R³⁶ is C₁-C₃ alkyl.

5 197. The method of Claim 196 wherein R³⁶ is methyl.

198. A method for the preparation of an alpha-amino acid compound having the structure of Formula 32:

10

32

or a salt, an enantiomer, or a racemate thereof,

wherein:

15

R² is selected from the group consisting of C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₅ alkoxy-C₁ alkyl, and C₁-C₅ alkylthio-C₁ alkyl;

R¹, R⁵, R⁶, and R⁷ independently are selected from the group consisting of -H, halogen, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, and C₁-C₅ alkoxy-C₁ alkyl;

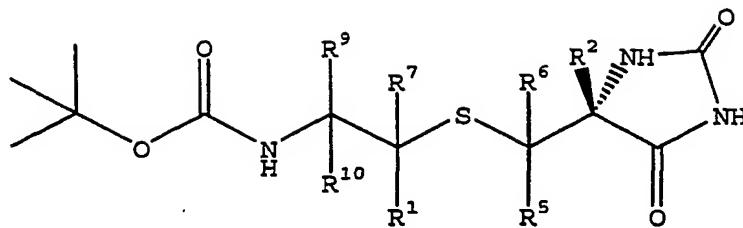
R⁹ and R¹⁰ independently are selected from the group consisting of -H, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, and C₁-C₅ alkoxy-C₁ alkyl; and

20

when any of R¹, R², R⁵, R⁶, R⁷, R⁹, and R¹⁰ independently is a moiety selected from the group consisting of alkyl, alkenyl, and alkynyl, then the moiety is optionally substituted by one or more substituent selected from the group consisting of -OH, alkoxy, and halogen;

25

wherein the method comprises treating under hydrolyzing conditions a hydantoin compound having the structure of Formula 33:

33

or a salt, an enantiomer, or a racemate thereof,
thereby forming the alpha-amino acid compound.

5

199. The method of Claim 198 wherein the hydrolyzing conditions comprise contacting the hydantoin compound with an acid to produce an acid hydrolyzate.

10 200. The method of Claim 199 wherein the acid is selected from the group consisting of hydrochloric acid, hydrobromic acid, hydroiodic acid, sulfuric acid, trifluoroacetic acid, and phosphoric acid.

201. The method of Claim 199 wherein the method further comprises treating the acid hydrolyzate with an ion exchange resin.

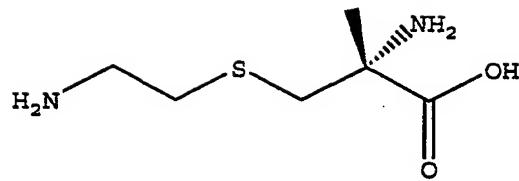
15 202. The method of Claim 198 wherein the hydrolyzing conditions comprise contacting the hydantoin compound with a base to produce a base hydrolyzate.

203. The method of Claim 202 wherein the base is selected from the group consisting of a hydrazine, a metal sulfide, a metal hydroxide, and a metal alkoxide

204. The method of Claim 198 wherein R¹, R⁵, R⁶, and R⁷ each is -H.

205. The method of Claim 198 wherein R⁹ and R¹⁰ each is -H.

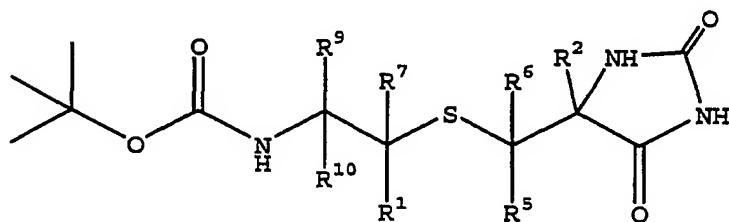
25 206. The method of Claim 198 where the alpha-amino acid compound has the structure of Formula 34:

34

or a salt, an enantiomer, or a racemate thereof.

5

207. A method for the preparation of a hydantoin compound having the structure of **Formula 35:**

35

10

or a salt, an enantiomer, or a racemate thereof,

wherein:

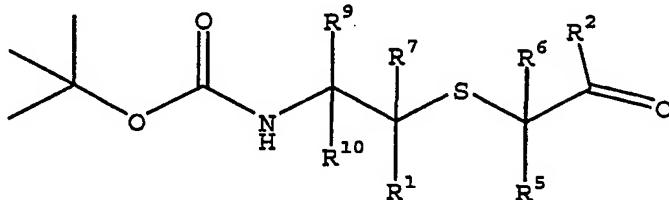
R² is selected from the group consisting of C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₅ alkoxy-C₁ alkyl, and C₁-C₅ alkylthio-C₁ alkyl;

R¹, R⁵, R⁶, and R⁷ independently are selected from the group consisting of -H, halogen, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, and C₁-C₅ alkoxy-C₁ alkyl;

R⁹ and **R¹⁰** independently are selected from the group consisting of -H, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, and C₁-C₅ alkoxy-C₁ alkyl; and

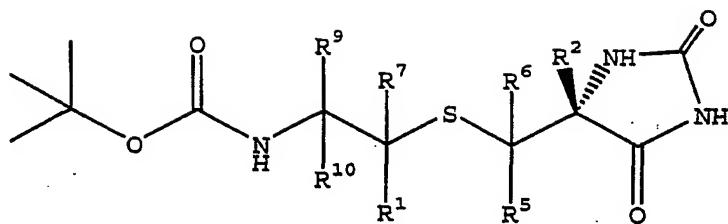
when any of **R¹, R², R⁵, R⁶, R⁷, R⁹, and R¹⁰** independently is a moiety selected from the group consisting of alkyl, alkenyl, and alkynyl, then the moiety is optionally substituted by one or more substituent selected from the group consisting of -OH, alkoxy, and halogen;

wherein the method comprises contacting an alpha-sulfo ketone compound having the structure of Formula 36:

36

5 with a source of cyanide in the presence of a source of ammonium carbonate and water, thereby producing the hydantoin compound.

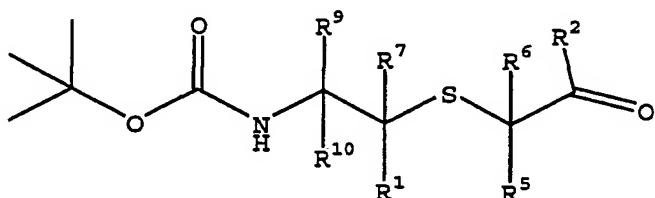
208. The method of Claim 207 wherein the source of cyanide is hydrogen cyanide.
- 10 209. The method of Claim 207 wherein the source of cyanide is a metal cyanide salt.
210. The method of Claim 209 wherein the metal cyanide salt is selected from the group consisting of sodium cyanide, potassium cyanide, and lithium cyanide.
- 15 211. The method of Claim 210 wherein the metal cyanide salt is sodium cyanide.
212. The method of Claim 207 wherein R¹, R⁵, R⁶, and R⁷ each is -H.
213. The method of Claim 207 wherein R⁹ and R¹⁰ each is -H.
- 20 214. The method of Claim 207 further comprising a chiral separation step.
215. The method of Claim 214 wherein the hydantoin compound has the structure of Formula 33:



33

or a salt or an enantiomer thereof.

5 216. A method for the preparation of an alpha-sulfo ketone compound having the structure of Formula 36:



36

wherein:

10 R^2 is selected from the group consisting of C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_1 - C_5 alkoxy- C_1 alkyl, and C_1 - C_5 alkylthio- C_1 alkyl;

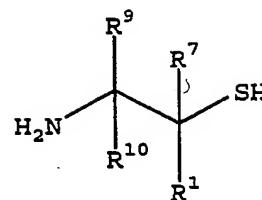
R^1 , R^5 , R^6 , and R^7 independently are selected from the group consisting of -H, halogen, C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, and C_1 - C_5 alkoxy- C_1 alkyl;

R^9 and R^{10} independently are selected from the group consisting of -H, C_1 - C_6

15 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, and C_1 - C_5 alkoxy- C_1 alkyl; and

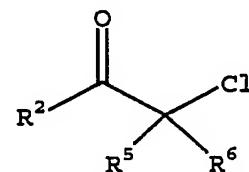
when any of R^1 , R^2 , R^5 , R^6 , R^7 , R^9 , and R^{10} independently is a moiety selected from the group consisting of alkyl, alkenyl, and alkynyl, then the moiety is optionally substituted by one or more substituent selected from the group consisting of -OH, alkoxy, and halogen;

20 wherein the method comprises contacting an aminothiol compound having the structure of Formula 37:

37

with di-*t*-butyl carbonate in the presence of a base to produce an intermediate mixture; and

5 contacting the intermediate mixture with an alpha-chloro ketone compound having the structure of Formula 38:

38

10 thereby producing the alpha-sulfo ketone compound.

217. The method of Claim 216 wherein the base is selected from the group consisting of a hydrazine, a metal sulfide, a metal hydroxide, a metal alkoxide, an amine, a hydroxylamine, and a metal amide complex.

15

218. The method of Claim 217 wherein the base is a metal hydroxide.

219. The method of Claim 218 wherein the base is selected from the group consisting of sodium hydroxide, potassium hydroxide, and lithium hydroxide.

20

220. The method of Claim 216 wherein R¹, R⁵, R⁶, and R⁷ each is -H.

221. The method of Claim 216 wherein R⁹ and R¹⁰ each is -H.

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